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| (54) Title: (R)-CHIRAL HALOGENATED 1-SUBSTITUTEDAMINO-(n+1)-ALKANOLS USEFUL FOR INHIBITING CHOLESTERYL ESTER TRANSFER PROTEIN ACTIVITY (57) Abstract The invention relates to substituted aryl and heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanol compounds useful as inhibitors of cholesteryl ester transfer protein (CETP; plasma lipid transfer protein-I) and compounds, compositions and methods for treating atherosclerosis and other coronary artery diseases. Novel high yield, stereoselective processes for the preparation of the chiral substituted alkanol compounds from chiral and achiral intermediates are described. | | |

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**(R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
Useful for Inhibiting Cholesteryl Ester Transfer Protein Activity**

FIELD OF THE INVENTION

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This invention is in the field of treating cardiovascular disease, and specifically relates to compounds, compositions, methods for treating atherosclerosis and other coronary artery disease, and methods for making compounds of this invention. More particularly, the invention relates to (R)-
10 chiral halogenated 1-substitutedamino-(n+1)-alkanol compounds that inhibit cholesteryl ester transfer protein (CETP), also known as plasma lipid transfer protein-I.

BACKGROUND OF THE INVENTION

15

Numerous studies have demonstrated that a low plasma concentration of high density lipoprotein (HDL) cholesterol is a powerful risk factor for the development of atherosclerosis (Barter and Rye, *Atherosclerosis*, 121, 1-12 (1996)). HDL is one of the major classes of lipoproteins that function in the
20 transport of lipids through the blood. The major lipids found associated with HDL include cholesterol, cholesteryl ester, triglycerides, phospholipids and fatty acids. The other classes of lipoproteins found in the blood are low density lipoprotein (LDL) and very low density lipoprotein (VLDL). Since low levels of HDL cholesterol increase the risk of atherosclerosis, methods for
25 elevating plasma HDL cholesterol would be therapeutically beneficial for the treatment of atherosclerosis and other diseases associated with accumulation of lipid in the blood vessels. These diseases include, but are not limited to, coronary heart disease, peripheral vascular disease, and stroke.

Atherosclerosis underlies most coronary artery disease (CAD), a major
30 cause of morbidity and mortality in modern society. High LDL cholesterol (above 180 mg/dl) and low HDL cholesterol (below 35 mg/dl) have been shown to be important contributors to the development of atherosclerosis. Other diseases, such as peripheral vascular disease, stroke, and hypercholesterolaemia are negatively affected by adverse HDL/LDL ratios.
35 Inhibition of CETP by the subject compounds is shown to effectively modify plasma HDL/LDL ratios, and to check the progress and/or formation of these diseases.

CETP is a plasma protein that facilitates the movement of cholesteryl esters and triglycerides between the various lipoproteins in the blood (Tall, *J. Lipid Res.*, 34, 1255-74 (1993)). The movement of cholesteryl ester from HDL to LDL by CETP has the effect of lowering HDL cholesterol. It therefore follows that inhibition of CETP should lead to elevation of plasma HDL cholesterol and lowering of plasma LDL cholesterol, thereby providing a therapeutically beneficial plasma lipid profile (McCarthy, *Medicinal Res. Revs.*, 13, 139-59 (1993); Sitori, *Pharmac. Ther.*, 67,443-47 (1995)). This exact phenomenon was first demonstrated by Swenson et al., (*J. Biol. Chem.*, 264, 14318 (1989)) with the use of a monoclonal antibody that specifically inhibited CETP. In rabbits, the antibody caused an elevation of the plasma HDL cholesterol and a decrease in LDL cholesterol. Son et al. (*Biochim. Biophys. Acta* 795, 743-480 (1984)), Morton et al. (*J. Lipid Res.* 35, 836-847 (1994)) and Tollefson et al. (*Am. J. Physiol.*, 255, (Endocrinol. Metab. 18, E894-E902 (1988))) describe proteins from human plasma that inhibit CETP. U.S. Patent 5,519,001, issued to Kushwaha et al., describes a 36 amino acid peptide derived from baboon apo C-I that inhibits CETP activity. Cho et al. (*Biochim. Biophys. Acta* 1391, 133-144 (1998)) describe a peptide from hog plasma that inhibits human CETP. Bonin et al. (*J. Peptide Res.*, 51, 216-225 (1998)) disclose a decapeptide inhibitor of CETP. A depsipeptide fungal metabolite is disclosed as a CETP inhibitor by Hedge et al. in *Bioorg. Med. Chem. Lett.*, 8, 1277-80 (1998).

There have been several reports of non-peptidic compounds that act as CETP inhibitors. Barrett et al. (*J. Am. Chem. Soc.*, 118, 7863-63 (1996)) and Kuo et al. (*J. Am. Chem. Soc.*, 117, 10629-34 (1995)) describe cyclopropane-containing CETP inhibitors. Pietzonka et al. (*Bioorg. Med. Chem. Lett.*, 6, 1951-54 (1996)) describe phosphonate-containing analogs of cholesteryl ester as CETP inhibitors. Coval et al. (*Bioorg. Med. Chem. Lett.*, 5, 605-610 (1995)) describe Wiedendiol-A and -B, and related sesquiterpene compounds as CETP inhibitors. Japanese Patent Application No. 10287662-A describes polycyclic, non-amine containing, polyhydroxylic natural compounds possessing CETP inhibition properties. Lee et al. (*J. Antibiotics*, 49, 693-96 (1996)) describe CETP inhibitors derived from an insect fungus. Busch et al. (*Lipids*, 25, 216-220, (1990)) describe cholesteryl acetyl bromide as a CETP inhibitor. Morton and Zilversmit (*J. Lipid Res.*, 35, 836-47 (1992)) describe that p-chloromercuriphenyl sulfonate, p-hydroxymercuribenzoate and ethyl mercurithiosalicylate inhibit CETP.

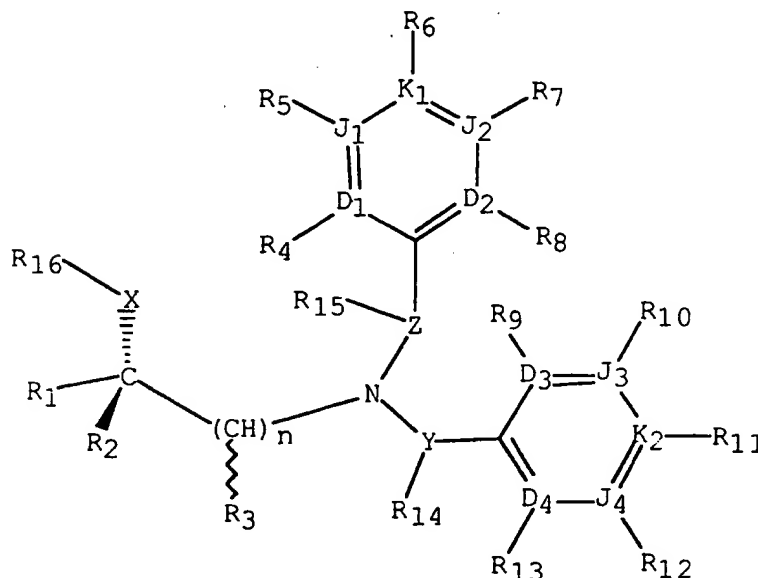
Connolly et al. (*Biochem. Biophys. Res. Comm.* 223, 42-47 (1996)) describe other cysteine modification reagents as CETP inhibitors. Xia et al. describe 1,3,5-triazines as CETP inhibitors (*Bioorg. Med. Chem. Lett.*, 6, 919-22 (1996)). Bisgaier et al. (*Lipids*, 29, 811-8 (1994)) describe 4-phenyl-5-
5 tridecyl-4H-1,2,4-triazole-thiol as a CETP inhibitor. Oomura et al. disclose non-peptidic tetracyclic and hexacyclic phenols as CETP inhibitors in Japanese Patent Application No. 10287662.

Some substituted heteroalkylamine compounds are known. In European Patent Application No. 796846, Schmidt et al. describe 2-aryl-
10 substituted pyridines as cholesteryl ester transfer protein inhibitors useful as cardiovascular agents. One substituent at C3 of the pyridine ring can be an hydroxyalkyl group. In European Patent Application No. 801060, Dow and Wright describe heterocyclic derivatives substituted with an aldehyde addition product of an alkylamine to afford 1-hydroxy-1-amines. These are reported to
15 be β 3-adrenergic receptor agonists useful for treating diabetes and other disorders. In Great Britain Patent Application No. 2305665, Fisher et al. disclose 3-agonist secondary amino alcohol substituted pyridine derivatives useful for treating several disorders including cholesterol levels and arteriosclerotic diseases. In European Patent Application No. 818448,
20 Schmidt et al. describe tetrahydroquinoline derivatives as cholesteryl ester transfer protein inhibitors. European Patent Application No. 818197, Schmek et al. describe pyridines with fused heterocycles as cholesteryl ester transfer protein inhibitors. Brandes et al. in German Patent Application No. 19627430 describe bicyclic condensed pyridine derivatives as cholesteryl ester transfer
25 protein inhibitors. In WO Patent Application No. 09839299, Muller-Gliemann et al. describe quinoline derivatives as cholesteryl ester transfer protein inhibitors. U.S. Patent 2,700,686, issued to Dickey and Towne, describes N-(2-haloalkyl-2-hydroxyethyl)amines in which the amine is further substituted with either 1 to 2 aliphatic groups or one aromatic group and one aliphatic
30 group. U.S. Patent 2,700,686 further describes a process to prepare the N-(2-haloalkyl-2-hydroxyethyl)amines by reacting halogenated-1,2-epoxyalkanes with the corresponding aliphatic amines and N-alkylanilines and their use as dye intermediates.

SUMMARY OF THE INVENTION

The present invention provides chiral compounds that can be used to inhibit cholesteryl ester transfer protein (CETP) activity and that have the

5 general structure:



In another aspect, the present invention includes pharmaceutical compositions comprising a pharmaceutically effective amount of the chiral compounds of this invention and a pharmaceutically acceptable carrier.

In another aspect, this invention relates to methods of using these chiral inhibitors as therapeutic agents in humans to inhibit cholesteryl ester transfer protein (CETP) activity, thereby decreasing the concentrations of low density lipoprotein (LDL) and raising the level of high density lipoprotein (HDL), resulting in a therapeutically beneficial plasma lipid profile. The compounds and methods of this invention can also be used to treat dyslipidemia (hypoalphalipoproteinemia), hyperlipoproteinaemia (chylomicronemia and hyperapobetalipoproteinemia), peripheral vascular disease, hypercholesterolaemia, atherosclerosis, coronary artery disease and other CETP-mediated disorders. The compounds can also be used in prophylactic treatment of subjects who are at risk of developing such disorders. The compounds can be used to lower the risk of atherosclerosis. The compounds of this invention would be also useful in prevention of cerebral vascular accident (CVA) or stroke. Besides being useful for human treatment, these

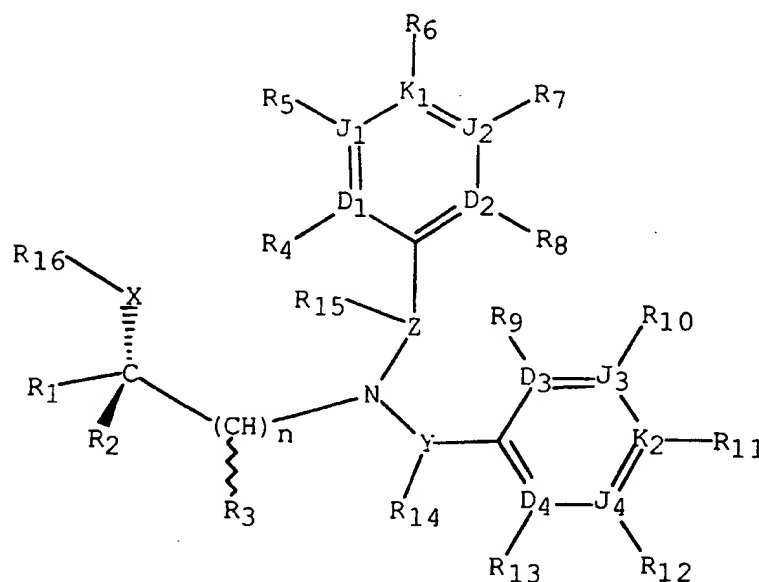
compounds are also useful for veterinary treatment of companion animals, exotic animals and farm animals such as primates, rabbits, pigs, horses, and the like.

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DESCRIPTION OF THE INVENTION

The present invention relates to a class of compounds comprising (R)-chiral halogenated 1-substitutedamino-(n+1)-alkanols which are beneficial in the therapeutic and prophylactic treatment of coronary artery disease as given in Formula I-H (also referred to herein as generic polycyclic aryl and heteroaryl

10 (R)-chiral halogenated 1-substitutedamino-(n+1)-alkanols):



(I-H)

or a pharmaceutically-acceptable salt thereof, wherein;

15 n is an integer selected from 1 through 4;

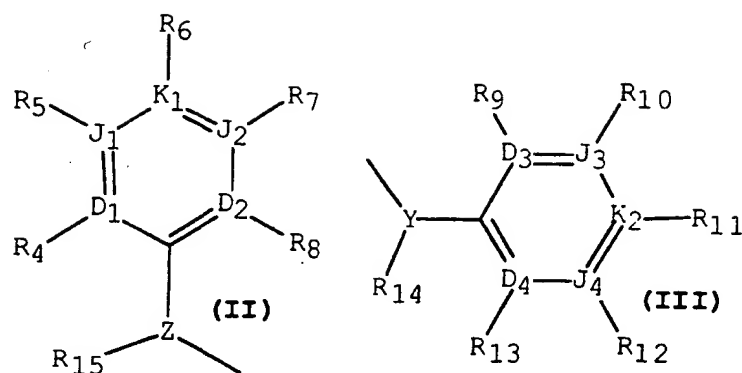
X is oxy;

R₁ is selected from the group consisting of haloalkyl, haloalkenyl,

haloalkoxymethyl, and haloalkenyloxymethyl with the proviso that R₁ has a

higher Cahn-Ingold-Prelog stereochemical system ranking than both R₂ and

20 (CHR₃)_n-N(A)Q wherein A is Formula (II) and Q is Formula (III);



- R_{16} is selected from the group consisting of hydrido, alkyl, alkenyl, alkynyl, aryl, aralkyl, aryloxyalkyl, alkoxyalkyl, alkenyloxyalkyl, alkylthioalkyl, arylthioalkyl, aralkoxyalkyl, heteroaralkoxyalkyl, alkylsulfinylalkyl, alkylsulfonylalkyl, cycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkenyl, cycloalkenylalkyl, haloalkyl, haloalkenyl, halocycloalkyl, halocycloalkenyl, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxyalkyl, halocycloalkenyloxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, heteroarylalkyl, monocarboalkoxyalkyl, monocarboalkoxy, dicarboalkoxyalkyl, monocarboxamido, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, acyl, aroyl, heteroaroyl, heteroaryloxyalkyl, dialkoxyposphonoalkyl, trialkylsilyl, and a spacer selected from the group consisting of a covalent single bond and a linear spacer moiety having a chain length of 1 to 4 atoms linked to the point of bonding of any aromatic substituent selected from the group consisting of R_4 , R_8 , R_9 , R_{13} , R_{14} , and R_{15} to form a heterocyclyl ring having from 5 through 10 contiguous members;

- D_1 , D_2 , J_1 , J_2 and K_1 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more than one of D_1 , D_2 , J_1 , J_2 and K_1 can be a covalent bond, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 can be O, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 can be S, one of D_1 , D_2 , J_1 , J_2 and K_1 must be a covalent bond when two

of D_1 , D_2 , J_1 , J_2 and K_1 are O and S, and no more than four of D_1 , D_2 , J_1 , J_2 and K_1 can be N;

D_3 , D_4 , J_3 , J_4 and K_2 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more
 5 than one can be a covalent bond, no more than one of D_3 , D_4 , J_3 , J_4 and K_2 can be O, no more than one of D_3 , D_4 , J_3 , J_4 and K_2 can be S, no more than two of D_3 , D_4 , J_3 , J_4 and K_2 can be O and S, one of D_3 , D_4 , J_3 , J_4 and K_2 must be a covalent bond when two of D_3 , D_4 , J_3 , J_4 and K_2 are O and S, and no more than four of D_3 , D_4 , J_3 , J_4 and K_2 can be N;

10 R_2 is hydrido;

R_2 can be selected from the group consisting of hydroxyalkyl, alkyl, alkenyl, alkynyl, aryl, aralkyl, aralkoxyalkyl, aryloxyalkyl, alkoxyalkyl, heteroaryloxyalkyl, alkenyloxyalkyl, cycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkenyl, cycloalkenylalkyl, haloalkyl, haloalkenyl,
 15 halocycloalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxyalkyl, heteroaryl, heteroarylalkyl, heteroarylthioalkyl, perhaloaryl, perhaloaralkyl, perhaloaralkyl, heteroaralkylthioalkyl, monocarboalkoxyalkyl, dicarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl,
 20 alkylsulfinylalkyl, alkylsulfonylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, cycloalkylsulfinylalkyl, cycloalkylsulfonylalkyl, heteroarylsulfonylalkyl, heteroarylsulfinylalkyl, aralkylsulfinylalkyl, aralkylsulfonylalkyl, carboxy, carboxyalkyl, carboalkoxy, carboxamide, carboxamidoalkyl, carboaralkoxy, dicyanoalkyl, carboalkoxycyanoalkyl, dialkoxyphosphonoalkyl, and
 25 diaralkoxyphosphonoalkyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$;

R_3 is selected from the group consisting of hydrido, hydroxy, halo, cyano, aryloxy, hydroxyalkyl, amino, alkylamino, dialkylamino, acyl,

- acylamido, alkoxy, alkyl, alkenyl, alkynyl, aryl, aralkyl, aryloxyalkyl, alkoxyalkyl, aralkoxyalkyl, alkylsulfinylalkyl, alkylsulfonylalkyl, aroyl, heteroaroyl, aralkylthioalkyl, heteroaralkylthioalkyl, heteroaryloxyalkyl, alkenyloxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl,
- 5 haloalkenyloxyalkyl, alkylthioalkyl, arylthioalkyl, cycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkenyl, cycloalkenylalkyl, heteroaryl, heteroarylalkyl, heteroarylthioalkyl, monocarboalkoxyalkyl, dicarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, arylsulfinylalkyl, arylsulfonylalkyl, cycloalkylsulfinylalkyl, cycloalkylsulfonylalkyl,
- 10 heteroarylsulfonylalkyl, heteroarylsulfinylalkyl, aralkylsulfinylalkyl, aralkylsulfonylalkyl, carboxy, carboxyalkyl, carboalkoxy, carboxamide, carboxamidoalkyl, carboaralkoxy, dialkoxyposphonoalkyl, and diaralkoxyposphonoalkyl with the provisos that $(\text{CHR}_3)_n\text{-N(A)Q}$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-
- 15 Ingold-Prelog stereochemical system ranking than R_2 ;
- Y is selected from a group consisting of a covalent single bond, $(\text{C}(\text{R}_{14})_2)_q$ wherein q is an integer selected from 1 through 2 and $(\text{CH}(\text{R}_{14}))_g\text{-W-(CH}(\text{R}_{14}))_p$ wherein g and p are integers independently selected from 0 through 1;
- 20 R_{14} is independently selected from the group consisting of hydrido, hydroxy, halo, cyano, aryloxy, amino, alkylamino, dialkylamino, hydroxyalkyl, acyl, aroyl, heteroaroyl, heteroaryloxyalkyl, sulfhydryl, acylamido, alkoxy, alkylthio, arylthio, alkyl, alkenyl, alkynyl, aryl, aralkyl, aryloxyalkyl, aralkoxyalkylalkoxy, alkylsulfinylalkyl, alkylsulfonylalkyl,
- 25 aralkylthioalkyl, heteroaralkoxythioalkyl, alkoxyalkyl, heteroaryloxyalkyl, alkenyloxyalkyl, alkylthioalkyl, arylthioalkyl, cycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkenyl, cycloalkenylalkyl, haloalkyl, haloalkenyl, halocycloalkyl, halocycloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl,
- 30 halocycloalkenyloxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, heteroarylalkyl, heteroarylthioalkyl, heteroaralkylthioalkyl, monocarboalkoxyalkyl, dicarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, alkylsulfinyl, alkylsulfonyl, haloalkylsulfinyl,

haloalkylsulfonyl, arylsulfinyl, arylsulfinylalkyl, arylsulfonyl,
arylsulfonylalkyl, aralkylsulfinyl, aralkylsulfonyl, cycloalkylsulfinyl,
cycloalkylsulfonyl, cycloalkylsulfinylalkyl, cycloalkylsulfonylalkyl,
heteroarylsulfonylalkyl, heteroarylsulfinyl, heteroarylsulfonyl,
5 heteroarylsulfinylalkyl, aralkylsulfinylalkyl, aralkylsulfonylalkyl, carboxy,
carboxyalkyl, carboalkoxy, carboxamide, carboxamidoalkyl, carboaralkoxy,
dialkoxyposphono, diaralkoxyposphono, dialkoxyposphonoalkyl,
diaralkoxyposphonoalkyl, a spacer selected from a moiety having a chain
length of 3 to 6 atoms connected to the point of bonding selected from the
10 group consisting of R₉ and R₁₃ to form a ring selected from the group
consisting of a cycloalkenyl ring having from 5 through 8 contiguous members
and a heterocyclyl ring having from 5 through 8 contiguous members, and a
spacer selected from a moiety having a chain length of 2 to 5 atoms connected
to the point of bonding selected from the group consisting of R₄ and R₈ to
15 form a heterocyclyl having from 5 through 8 contiguous members with the
proviso that, when Y is a covalent bond, an R₁₄ substituent is not attached to
Y;

R₁₄ and R₁₅ can be taken together to form a spacer selected from a
moiety having a chain length of 2 to 5 atoms to form a heterocyclyl ring having
20 from 5 through 8 contiguous members;

R₁₄ and R₁₄, when bonded to the different atoms, can be taken
together to form a group selected from the group consisting of a covalent bond,
alkylene, haloalkylene, and a spacer selected from a group consisting of a
moiety having a chain length of 2 to 5 atoms connected to form a ring selected
25 from the group of a saturated cycloalkyl having from 5 through 8 contiguous
members, a cycloalkenyl having from 5 through 8 contiguous members, and a
heterocyclyl having from 5 through 8 contiguous members;

R₁₄ and R₁₄, when bonded to the same atom can be taken together to
form a group selected from the group consisting of oxo, thiono, alkylene,
30 haloalkylene, and a spacer selected from the group consisting of a moiety
having a chain length of 3 to 7 atoms connected to form a ring selected from the
group consisting of a cycloalkyl having from 4 through 8 contiguous

members, a cycloalkenyl having from 4 through 8 contiguous members, and a heterocyclyl having from 4 through 8 contiguous members:

W is selected from the group consisting of O, C(O), C(S),

C(O)N(R₁₄), C(S)N(R₁₄), (R₁₄)NC(O), (R₁₄)NC(S), S, S(O), S(O)₂,

- 5 S(O)₂N(R₁₄), (R₁₄)NS(O)₂, and N(R₁₄) with the proviso that R₁₄ is selected from other than halo and cyano;

Z is independently selected from a group consisting of a covalent single bond, (C(R₁₅)₂)_q wherein q is an integer selected from 1 through 2.

(CH(R₁₅))_j-W-(CH(R₁₅))_k wherein j and k are integers independently

- 10 selected from 0 through 1 with the proviso that, when Z is a covalent single bond, an R₁₅ substituent is not attached to Z;

- R₁₅ is independently selected, when Z is (C(R₁₅)₂)_q wherein q is an integer selected from 1 through 2, from the group consisting of hydrido, hydroxy, halo, cyano, aryloxy, amino, alkylamino, dialkylamino, hydroxyalkyl, acyl, aroyl, heteroaroyl, heteroaryloxyalkyl, sulfhydryl, acylamido, alkoxy, alkylthio, arylthio, alkyl, alkenyl, alkynyl, aryl, aralkyl, aryloxyalkyl, aralkoxyalkyl, alkylsulfanylalkyl, alkylsulfonylalkyl, aralkylthioalkyl, heteroaralkylthioalkyl, alkoxyalkyl, heteroaryloxyalkyl, alkenyloxyalkyl, alkylthioalkyl, arylthioalkyl, cycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkenyl, cycloalkenylalkyl, haloalkyl, haloalkenyl, halocycloalkyl, halocycloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, heteroarylalkyl, heteroarylthioalkyl, heteroaralkylthioalkyl, monocarboalkoxyalkyl, dicarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, alkylsulfinyl, alkylsulfonyl, haloalkylsulfinyl, haloalkylsulfonyl, arylsulfinyl, arylsulfinylalkyl, arylsulfonyl, arylsulfonylalkyl, aralkylsulfinyl, aralkylsulfonyl, cycloalkylsulfinyl, cycloalkylsulfonyl, cycloalkylsulfinylalkyl, cycloalkylsulfonylalkyl, heteroarylsulfonylalkyl, heteroarylsulfinyl, heteroarylsulfonyl, heteroarylsulfinylalkyl, aralkylsulfinylalkyl, aralkylsulfonylalkyl, carboxy, carboxyalkyl, carboalkoxy, carboxamide, carboxamidoalkyl, carboaralkoxy, dialkoxyposphono, diaralkoxyposphono, dialkoxyposphonoalkyl.
- 15
- 20
- 25
- 30

diaralkoxyphosphonoalkyl, a spacer selected from a moiety having a chain length of 3 to 6 atoms connected to the point of bonding selected from the group consisting of R_4 and R_8 to form a ring selected from the group consisting of a cycloalkenyl ring having from 5 through 8 contiguous members and a heterocyclyl ring having from 5 through 8 contiguous members, and a spacer selected from a moiety having a chain length of 2 to 5 atoms connected to the point of bonding selected from the group consisting of R_9 and R_{13} to form a heterocyclyl having from 5 through 8 contiguous members;

R_{15} and R_{15} , when bonded to the different atoms, can be taken together to form a group selected from the group consisting of a covalent bond, alkylene, haloalkylene, and a spacer selected from a group consisting of a moiety having a chain length of 2 to 5 atoms connected to form a ring selected from the group of a saturated cycloalkyl having from 5 through 8 contiguous members, a cycloalkenyl having from 5 through 8 contiguous members, and a heterocyclyl having from 5 through 8 contiguous members;

R_{15} and R_{15} , when bonded to the same atom, can be taken together to form a group selected from the group consisting of oxo, thiono, alkylene, haloalkylene, and a spacer selected from the group consisting of a moiety having a chain length of 3 to 7 atoms connected to form a ring selected from the group consisting of a cycloalkyl having from 4 through 8 contiguous members, a cycloalkenyl having from 4 through 8 contiguous members, and a heterocyclyl having from 4 through 8 contiguous members;

R_{15} is independently selected, when Z is $(CH(R_{15}))_j-W-(CH(R_{15}))_k$ wherein j and k are integers independently selected from 0 through 1, from the group consisting of hydrido, halo, cyano, aryloxy, carboxyl, acyl, aroyl, heteroaroyl, hydroxyalkyl, heteroaryloxyalkyl, acylamido, alkoxy, alkylthio, arylthio, alkyl, alkenyl, alkynyl, aryl, aralkyl, aryloxyalkyl, alkoxyalkyl, heteroaryloxyalkyl, aralkoxyalkyl, heteroaralkoxyalkyl, alkylsulfonylalkyl, alkylsulfinylalkyl, alkenyloxyalkyl, alkylthioalkyl, arylthioalkyl, cycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkenyl, cycloalkenylalkyl, haloalkyl, haloalkenyl, halocycloalkyl, halocycloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl,

heteroaryl, heteroaralkyl, heteroarylthioalkyl, heteroaralkylthioalkyl, monocarboalkoxyalkyl, dicarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, alkylsulfinyl, alkylsulfonyl, haloalkylsulfinyl, haloalkylsulfonyl, arylsulfinyl, arylsulfinylalkyl, arylsulfonyl, arylsulfonylalkyl, aralkylsulfinyl, aralkylsulfonyl, cycloalkylsulfinyl, cycloalkylsulfonyl, cycloalkylsulfinylalkyl, cycloalkylsulfonylalkyl, heteroarylsulfonylalkyl, heteroarylsulfinyl, heteroarylsulfonyl, heteroarylsulfinylalkyl, aralkylsulfinylalkyl, aralkylsulfonylalkyl, carboxyalkyl, carboalkoxy, carboxamide, carboxamidoalkyl, carboaralkoxy, dialkoxyphosphonoalkyl, diaralkoxyphosphonoalkyl, a spacer selected from a linear moiety having a chain length of 3 to 6 atoms connected to the point of bonding selected from the group consisting of R₄ and R₈ to form a ring selected from the group consisting of a cycloalkenyl ring having from 5 through 8 contiguous members and a heterocyclyl ring having from 5 through 8 contiguous members, and a spacer selected from a linear moiety having a chain length of 2 to 5 atoms connected to the point of bonding selected from the group consisting of R₉ and R₁₃ to form a heterocyclyl ring having from 5 through 8 contiguous members;

R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁, R₁₂, and R₁₃ are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroarylamino, N-heteroarylamino-N-alkylamino, heteroaralkyl, heteroarylaminomethyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxylalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro, lower alkylamino, alkylthio, alkylthioalkyl, arylamino, aralkylamino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, heteroarylsulfinylalkyl, heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl,

- haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, heterocyclisulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl with the proviso that there are one to five non-hydrido ring substituents R₄, R₅, R₆, R₇, and R₈ present, that there are one to five non-hydrido ring substituents R₉, R₁₀, R₁₁, R₁₂, and R₁₃ present, and R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁, R₁₂, and R₁₃ are each independently selected to maintain the tetravalent nature of carbon, trivalent nature of nitrogen, the divalent nature of sulfur, and the divalent nature of oxygen;
- R₄ and R₅, R₅ and R₆, R₆ and R₇, R₇ and R₈, R₉ and R₁₀, R₁₀ and R₁₁, R₁₁ and R₁₂, and R₁₂ and R₁₃ can be independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no

more than one of the group consisting of spacer pairs R₄ and R₅, R₅ and R₆, R₆ and R₇, and R₇ and R₈, can be used at the same time and that no more than one of the group consisting of spacer pairs R₉ and R₁₀, R₁₀ and R₁₁, R₁₁ and R₁₂, and R₁₂ and R₁₃ can be used at the same time;

5 R₄ and R₉, R₄ and R₁₃, R₈ and R₉, and R₈ and R₁₃ can be independently selected to form a spacer pair wherein said spacer pair is taken together to form a linear moiety wherein said linear moiety forms a ring selected from the group consisting of a partially saturated heterocyclyl ring having from 5 through 8 contiguous members and a heteroaryl ring having
10 from 5 through 6 contiguous members with the proviso that no more than one of the group consisting of spacer pairs R₄ and R₉, R₄ and R₁₃, R₈ and R₉, and R₈ and R₁₃ can be used at the same time;

 R₅ and R₁₀, R₅ and R₁₂, R₇ and R₁₀, and R₇ and R₁₂ can be independently selected to form a spacer pair wherein said spacer pair is taken
15 together to form a linear moiety wherein said linear moiety forms a C8 to C13 heterocyclyl ring having from 8 through 13 contiguous members with the proviso that no more than one of the group consisting of spacer pairs R₅ and R₁₀, R₅ and R₁₂, R₇ and R₁₀, and R₇ and R₁₂ can be used at the same time.

20 In another embodiment of compounds of Formula I-H,

 D₁, D₂, J₁, J₂ and K₁ are each carbon with the proviso that at least one of D₃, D₄, J₃, J₄ and K₂ is selected from the group consisting of O, S, and N, wherein D₃, D₄, J₃, J₄ and K₂ are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no
25 more than one of D₃, D₄, J₃, J₄ and K₂ can be a covalent bond, no more than one of D₃, D₄, J₃, J₄ and K₂ can be O, no more than one of D₃, D₄, J₃, J₄

and K_2 can be S, one of D_3, D_4, J_3, J_4 and K_2 must be a covalent bond when two of D_3, D_4, J_3, J_4 and K_2 are O and S, and no more than four of D_3, D_4, J_3, J_4 and K_2 can be N;

D_1, D_2, J_1, J_2 and K_1 can be selected from the group consisting of C,

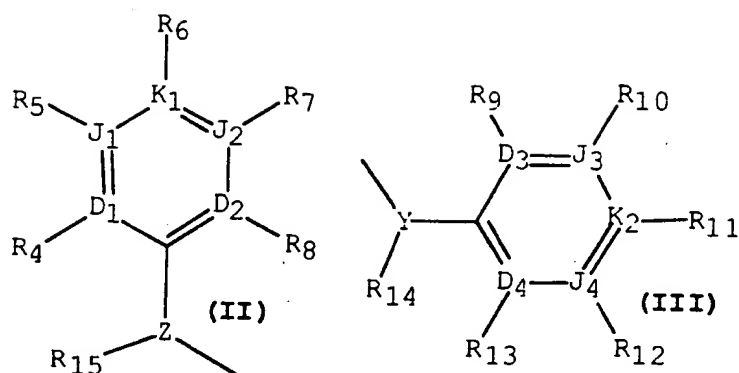
- 5 O, S, N and covalent bond with the provisos that D_3, D_4, J_3, J_4 and K_2 are each carbon and at least one of D_1, D_2, J_1, J_2 and K_1 is selected from the group consisting of O, S, and N wherein, when D_1, D_2, J_1, J_2 and K_1 are selected from the group consisting of C, O, S, covalent bond, and N, no more than one of D_1, D_2, J_1, J_2 and K_1 can be a covalent bond, no more than one
- 10 of D_1, D_2, J_1, J_2 and K_1 can be O, no more than one of D_1, D_2, J_1, J_2 and K_1 can be S, one of D_1, D_2, J_1, J_2 and K_1 must be a covalent bond when two of D_1, D_2, J_1, J_2 and K_1 are O and S, and no more than four of D_1, D_2, J_1, J_2 and K_1 can be N;

n is an integer selected from 1 through 4;

- 15 X is oxy;

R_{16} is selected from the group consisting of hydrido, acyl, aroyl, and trialkylsilyl;

- R_1 is selected from the group consisting of haloalkyl, haloalkenyl, haloalkoxymethyl, and haloalkenyloxymethyl with the proviso that R_1 has a
- 20 higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);



R_2 is hydrido;

R_2 can be selected from the group consisting of aryl, aralkyl, alkyl,

- 5 alkenyl, alkenyloxyalkyl, haloalkyl, haloalkenyl, halocycloalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, dicyanoalkyl, and carboalkoxycyanoalkyl with the proviso that R_2 has a lower Cahn-Ingold-

Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$;

- 10 R_3 is selected from the group consisting of hydrido, hydroxy, cyano, aryl, aralkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, heteroaryl, alkenyloxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-

- 15 Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from the group consisting of covalent single bond and $(C(R_{14})_2)_q$ wherein q is an integer selected from 1 through 2;

R_{14} is selected from the group consisting of hydrido, cyano,

- 20 hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl;

Z is selected from the group consisting of covalent single bond,

$(C(R_{15})_2)_q$ wherein q is an integer selected from 1 through 2, and

$(CH(R_{15}))_j-W-(CH(R_{15}))_k$ wherein j and k are integers independently selected from 0 through 1;

5 W is oxy;

R_{15} is selected from the group consisting of hydrido, cyano, hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl;

10 R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido, halo, haloalkyl, and alkyl ;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroarylsulfonyl, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, cycloalkoxy, cycloalkylalkoxy, hydroxy, amino, thio, nitro, lower alkylamino, alkylthio, arylamino, aralkylamino, arylthio, arylthioalkyl, alkylsulfonamido, monoarylamidosulfonyl, arylsulfonamido, heteroarylthio, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, halo, haloalkyl, haloalkoxy, hydroxyhaloalkyl, hydroxyalkyl, aryl, aryloxy, aralkoxy, saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, carboalkoxy, alkoxy-carboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboxamido, carboxamidoalkyl, and cyano;

R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} spacer pairs can be independently selected from the group consisting of alkylene, alkenylene, alkylenedioxy, aralkylene, diacyl, haloalkylene, and aryloxylylene with the provisos that no

more than one of the group consisting of spacer pairs R₄ and R₅, R₅ and R₆, R₆ and R₇, and R₇ and R₈ can be used at the same time and that no more than one of the group consisting of spacer pairs R₉ and R₁₀, R₁₀ and R₁₁, R₁₁ and R₁₂, and R₁₂ and R₁₃ can be used at the same time.

5

In an even more specific embodiment of compounds Formula I-H,

D₁, D₂, J₁, J₂ and K₁ are each carbon;

D₃, D₄, J₃, J₄ and K₂ are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that at least one of D₃, D₄, J₃, J₄ and K₂ is selected from the group consisting of O, S, and N, wherein no more than one of D₃, D₄, J₃, J₄ and K₂ can be a covalent bond, no more than one of D₃, D₄, J₃, J₄ and K₂ can be O, no more than one of D₃, D₄, J₃, J₄ and K₂ can be S, one of D₃, D₄, J₃, J₄ and K₂ must be a covalent bond when two of D₃, D₄, J₃, J₄ and K₂ are O and S, and no more than four of D₃, D₄, J₃, J₄ and K₂ can be N;

n is an integer selected from 1 to 3;

X is oxy;

R₁ is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, chloromethyl, fluoromethyl, difluoromethyl, chlorodifluoromethyl, pentafluoroethyl, 2,2,3,3,3-pentafluoropropyl, and heptafluoropropyl;

R₁₆ is selected from the group consisting of acetyl, benzoyl, dimethyl *tert*-butylsilyl, hydrido, and trimethylsilyl;

R₂ is hydrido;

R₂ can be selected from the group consisting of hydrido, methyl, ethyl, propyl, butyl, isopropyl, isobutyl, vinyl, phenyl,

4-trifluoromethylphenyl, 1,1,2,2-tetrafluoroethoxymethyl, chloromethyl, trifluoromethoxymethyl, fluoromethyl, difluoromethyl, 2,2,3,3,3-pentafluoropropyl, and pentafluorophenoxymethyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n$ -

5 N(A)Q:

R_3 is selected from the group consisting of hydrido, hydroxy, cyano, acetyl, methoxy, ethoxy, methyl, ethyl, propyl, vinyl, phenyl, methoxymethyl, 4-trifluoromethylphenyl, trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, chloromethyl, fluoromethyl, difluoromethyl, 10 chlorodifluoromethyl, pentafluoroethyl, 2,2,3,3,3-pentafluoropropyl, heptafluoropropyl, pentafluorophenyl, and pentafluorophenoxymethyl with the provisos that $(CHR_3)_n$ -N(A)Q has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 .

15

In another even more specific embodiment of compounds Formula I-H, D_3 , D_4 , J_3 , J_4 and K_2 are each carbon;

D_1 , D_2 , J_1 , J_2 and K_1 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that at least one 20 of D_1 , D_2 , J_1 , J_2 and K_1 is selected from the group consisting of O, S, and N, wherein no more than one of D_1 , D_2 , J_1 , J_2 and K_1 can be a covalent bond, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 can be O, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 can be S, one of D_1 , D_2 , J_1 , J_2 and K_1 must be a covalent bond when two of D_1 , D_2 , J_1 , J_2 and K_1 are O and S, and no more 25 than four of D_1 , D_2 , J_1 , J_2 and K_1 can be N;

n is an integer selected from 1 to 3;

X is oxy;

R_1 is selected from the group consisting of trifluoromethyl,

1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, chloromethyl, fluoromethyl, difluoromethyl, chlorodifluoromethyl, pentafluoroethyl, 2,2,3,3,3-pentafluoropropyl, and heptafluoropropyl;

- 5 R_{16} is selected from the group consisting of acetyl, benzoyl, dimethyl *tert*-butylsilyl, hydrido, and trimethylsilyl:

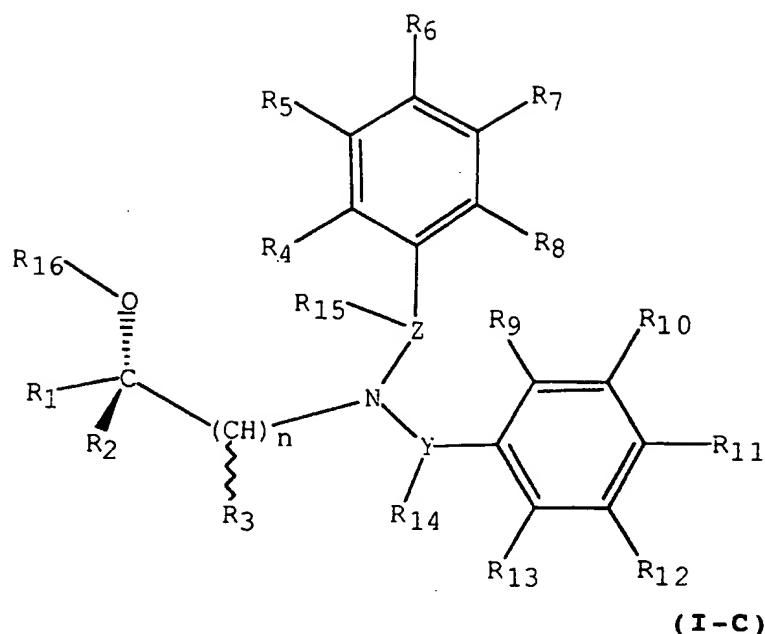
R_2 is hydrido;

- R_2 can be selected from the group consisting of hydrido, methyl, ethyl, propyl, butyl, isopropyl, isobutyl, vinyl, phenyl, 10 4-trifluoromethylphenyl, 1,1,2,2-tetrafluoroethoxymethyl, chloromethyl, trifluoromethoxymethyl, fluoromethyl, difluoromethyl, 2,2,3,3,3-pentafluoropropyl, and pentafluorophenoxymethyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$;

- 15 R_3 is selected from the group consisting of hydrido, hydroxy, cyano, acetyl, methoxy, ethoxy, methyl, ethyl, propyl, vinyl, phenyl, methoxymethyl, 4-trifluoromethylphenyl, trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, chloromethyl, fluoromethyl, difluoromethyl, chlorodifluoromethyl, pentafluoroethyl, 2,2,3,3,3-pentafluoropropyl, 20 heptafluoropropyl, pentafluorophenyl, and pentafluorophenoxymethyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 .

- 25 In a preferred embodiment of compounds of Formula I-H, the compounds correspond to the Formula I-C (also referred to herein as phenyl (R)-chiral halogenated 1-substitutedamino-(n+1)-alkanols):

21



or a pharmaceutically acceptable salt thereof, wherein:

n is an integer selected from 1 through 4;

5 R_{16} is selected from the group consisting of hydrido, alkyl, acyl,

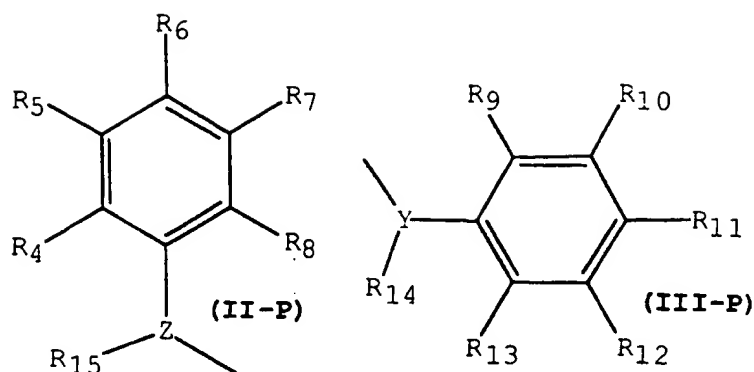
aroyl, heteroaroyl, trialkylsilyl, and a spacer selected from the group consisting of a covalent single bond and a linear spacer moiety having a chain length of 1 to 4 atoms linked to the point of bonding of any aromatic substituent selected from the group consisting of R_4 , R_8 , R_9 , and R_{13} to form a heterocyclyl ring

10 having from 5 through 10 contiguous members with the proviso that said linear spacer moiety is other than covalent single bond when R_2 is alkyl;

R_1 is selected from the group consisting of haloalkyl, haloalkenyl,

haloalkoxymethyl, and haloalkenyloxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and

15 $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



R_2 is hydrido;

R_2 can be selected from the group consisting of aryl, aralkyl, alkyl, alkenyl, alkenyloxyalkyl, haloalkyl, haloalkenyl, halocycloalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, dicyanoalkyl, and carboalkoxycyanoalkyl with the proviso that R_2 has a lower Cahn-Ingold-

Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

R_3 is selected from the group consisting of hydrido, hydroxy, cyano, aryl, aralkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, heteroaryl, alkenyloxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-

Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from the group consisting of covalent single bond and $(C(R_{14})_2)_q$ wherein q is an integer selected from 1 through 2;

R_{14} is selected from the group consisting of hydrido, hydroxy, cyano, hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkynyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, carboalkoxy, carboxamide, carboxamidoalkyl;

Z is selected from the group consisting of covalent single bond.

$(C(R_{15})_2)_q$ wherein q is an integer selected from 1 through 2, and

$(CH(R_{15}))_j-W-(CH(R_{15}))_k$ wherein j and k are integers independently selected from 0 through 1;

5 W is selected from the group consisting of O, C(O), C(S),

C(O)N(R₁₄), C(S)N(R₁₄), (R₁₄)NC(O), (R₁₄)NC(S), S, S(O), S(O)₂,

S(O)₂N(R₁₄), (R₁₄)NS(O)₂, and N(R₁₄) with the proviso that R₁₄ is other than cyano;

R₁₅ is selected from the group consisting of hydrido, cyano,

10 hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkynyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, carboalkoxy, carboxamide, and carboxamidoalkyl;

R₄, R₈, R₉, and R₁₃ are independently selected from the group

15 consisting of hydrido, halo, haloalkyl, and alkyl ;

R₅, R₆, R₇, R₁₀, R₁₁, and R₁₂ are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl, 20 aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroaryl amino, N-heteroaryl amino-N-alkyl amino, heteroaryl aminoalkyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxylalkyl, heteroaralkoxy, 25 cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro, lower alkyl amino, alkylthio, alkylthioalkyl, aryl amino, aralkyl amino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, 30 alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, heteroaryl sulfinylalkyl, heteroaryl sulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl,

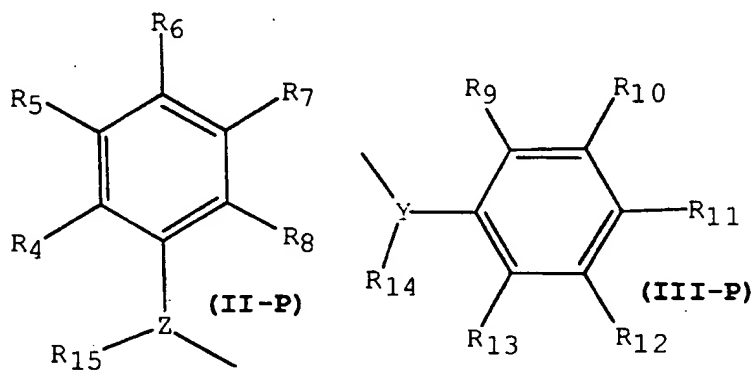
- alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl;

- R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} can be independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no more than one of the group consisting of spacer pairs R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , can be used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} can be used at the same time.

- In a preferred embodiment of compounds of Formula I-C, n is an integer selected from 1 through 4;

R_{16} is selected from the group consisting of hydrido, acyl, aroyl, and trialkylsilyl;

R_1 is selected from the group consisting of haloalkyl, haloalkenyl, haloalkoxymethyl, and haloalkenyloxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



R_2 is hydrido;

R_2 can be selected from the group consisting of aryl, aralkyl, alkyl, alkenyl, alkenyloxyalkyl, haloalkyl, haloalkenyl, halocycloalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, dicyanoalkyl, and carboalkoxycyanoalkyl with the proviso that R_2 has a lower Cahn-Ingold-

Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

R_3 is selected from the group consisting of hydrido, hydroxy, cyano, aryl, aralkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, heteroaryl, alkenyloxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-

Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-

Ingold-Prelog stereochemical system ranking than R_2 :

Y is selected from the group consisting of covalent single bond and $(C(R_{14})_2)_q$ wherein q is an integer selected from 1 through 2:

- 5 R_{14} is selected from the group consisting of hydrido, cyano, hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl:

Z is selected from the group consisting of covalent single bond,

- 10 $(C(R_{15})_2)_q$ wherein q is an integer selected from 1 through 2, and $(CH(R_{15}))_j-W-(CH(R_{15}))_k$ wherein j and k are integers independently selected from 0 through 1;

W is oxy;

- R_{15} is selected from the group consisting of hydrido, cyano, 15 hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and carboxamidoalkyl;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido, halo, haloalkyl, and alkyl ;

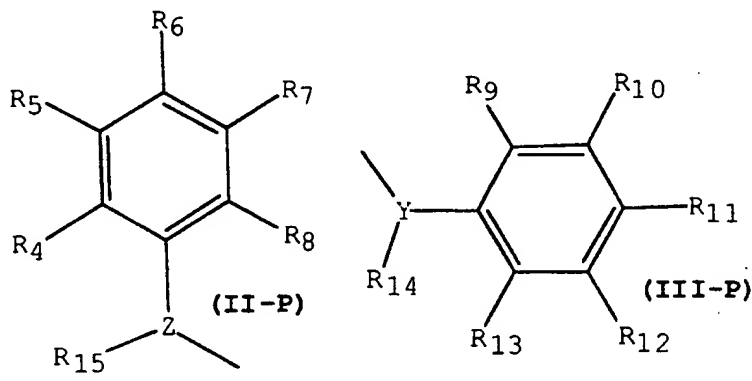
- 20 R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroarylsulfonyl, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, cycloalkoxy, 25 cycloalkylalkoxy, hydroxy, amino, thio, nitro, lower alkylamino, alkylthio, arylamino, aralkylamino, arylthio, arylthioalkyl, alkylsulfonyl, alkylsulfonamido, monoarylamidosulfonyl, arylsulfonyl, heteroarylthio, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkylenedioxy, haloalkylenedioxy, cycloalkyl, 30 cycloalkylalkanoyl, halo, haloalkyl, haloalkoxy, hydroxyhaloalkyl, hydroxyalkyl, aryl, aralkyl, aryloxy, aralkoxy, saturated heterocyclyl,

heteroaryl, heteroaryloxy, heteroaryloxyalkyl, arylalkyl, heteroarylalkyl, arylalkenyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboxamido, carboxamidoalkyl, and cyano;

- 5 R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} spacer pairs can be independently selected from the group consisting of alkylene, alkenylene, alkylenedioxy, aralkylene, diacyl, haloalkylene, and aryloxylylene with the provisos that no more than one of the group consisting of spacer pairs R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 can be used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} can be used at the same time.

In a more preferred embodiment of compounds of Formula I-C, n is an integer selected from 1 through 2;

- 15 R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



- 20 R_{16} is hydrido;

R_2 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, haloalkoxy, haloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, and heteroaryl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

- 5 R_3 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, and haloalkoxyalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

- 10 Y is selected from the group consisting of a covalent single bond and alkylene;

Z is selected from the group consisting of a covalent single bond and alkylene;

R_{14} is selected from the group consisting of hydrido, alkyl, and haloalkyl;

- 15 R_{15} is selected from the group consisting of hydrido, alkyl, and haloalkyl;

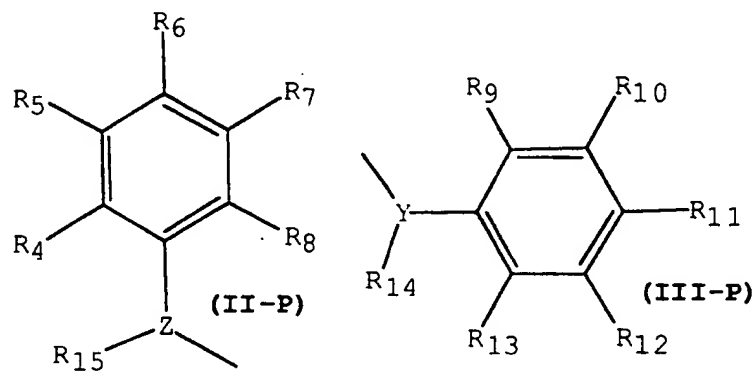
R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

- 20 R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

- 25 In an even more preferred embodiment of compounds of Formula I-C, n is the integer 1;

R_{16} is hydrido;

R_1 is haloalkyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



5 R_2 is hydrido;

R_2 can be selected from the group consisting of alkyl, haloalkyl, aryl, and haloalkoxy with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

R_3 is selected from the group consisting of hydrido, alkyl, and
10 haloalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is alkylene;

Z is covalent single bond;

15 R_{14} is hydrido;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio,
20 arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy,

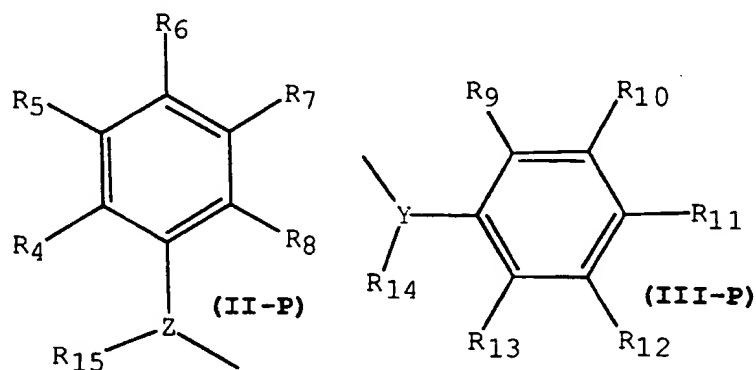
alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, and heteroaryloxyalkyl.

In an embodiment of compounds of Formula I-C,
 5 n is an integer selected from 1 to 3;

R_1 is selected from the group consisting of trifluoromethyl,

1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, chloromethyl, fluoromethyl, difluoromethyl, chlorodifluoromethyl, pentafluoroethyl, 2,2,3,3,3-pentafluoropropyl, and heptafluoropropyl with the

10 proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



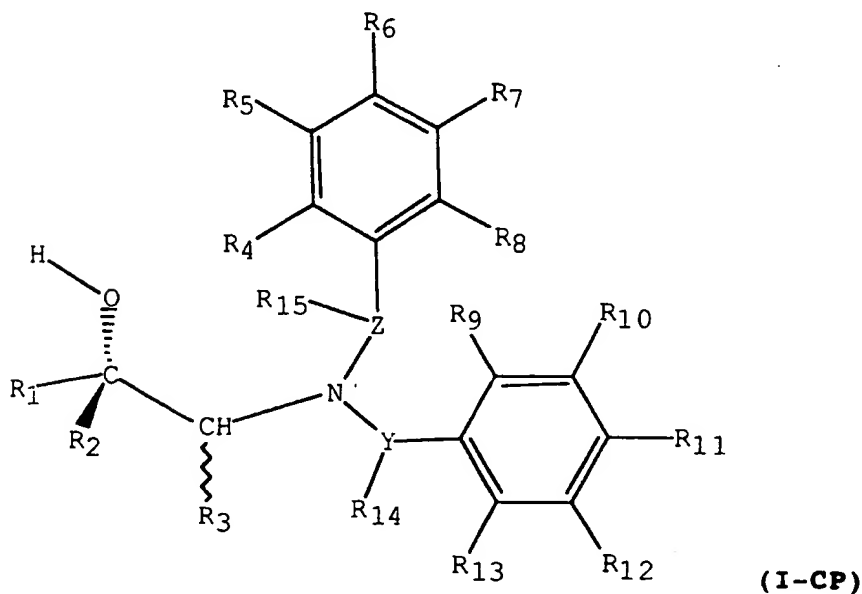
R_{16} is selected from the group consisting of acetyl, benzoyl, dimethyl
 15 *tert*-butylsilyl, hydrido, and trimethylsilyl;

R_2 is hydrido;

R_2 can be selected from the group consisting of methyl, ethyl, propyl, butyl, isopropyl, isobutyl, vinyl, phenyl, 4-trifluoromethylphenyl, 1,1,2,2-tetrafluoroethoxymethyl, chloromethyl, trifluoromethoxymethyl, fluoromethyl, difluoromethyl, 2,2,3,3,3-pentafluoropropyl, and and
 20 pentafluorophenoxymethyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

- R_3 is selected from the group consisting of hydrido, hydroxy, cyano, acetyl, methoxy, ethoxy, methyl, ethyl, propyl, vinyl, phenyl, methoxymethyl, 4-trifluoromethylphenyl, trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, chloromethyl, fluoromethyl, difluoromethyl, chlorodifluoromethyl, pentafluoroethyl, 2,2,3,3,3-pentafluoropropyl, heptafluoropropyl, pentafluorophenyl, and pentafluorophenoxymethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 .

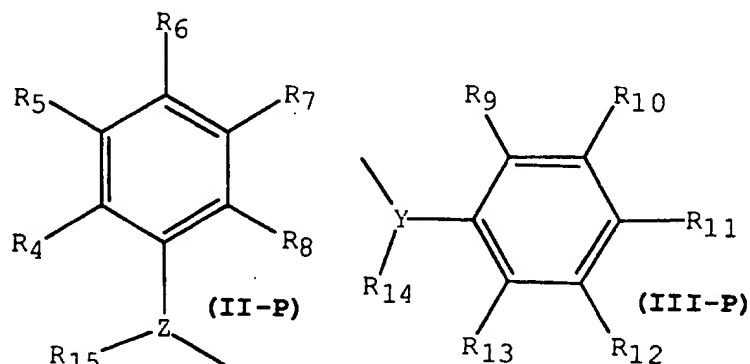
- 10 In a preferred embodiment of compounds of Formula I-C, compounds have the Formula I-CP:



or a pharmaceutically acceptable salt thereof, wherein;

- R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a

higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P):



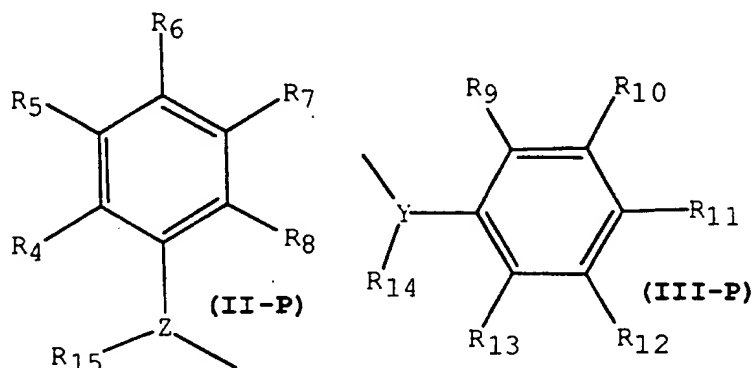
R_2 is hydrido:

- 5 R_2 can be selected from the group consisting of methyl, ethyl, propyl, butyl, vinyl, phenyl, 4-trifluoromethylphenyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:
- 10 R_3 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, methyl, ethyl, vinyl, methoxymethyl, trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-
- 15 Ingold-Prelog stereochemical system ranking than R_2 .

In a even more preferred embodiment of compounds of Formula I-CP,

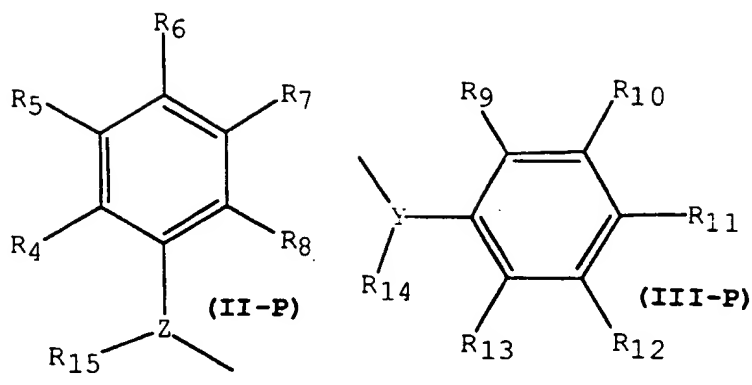
- R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl,
- 20 chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a

higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P):



R_2 is hydrido:

- 5 R_2 can be selected from the group consisting of methyl, ethyl, phenyl, 4-trifluoromethylphenyl, trifluoromethoxymethyl, 1,1,2,2-tetrafluoroethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:
- 10 R_3 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, methyl, trifluoromethyl, difluoromethyl, and chlorodifluoromethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 .
- 15 In a most preferred embodiment of compounds of Formula I-CP, R_1 is selected from the group consisting of trifluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);

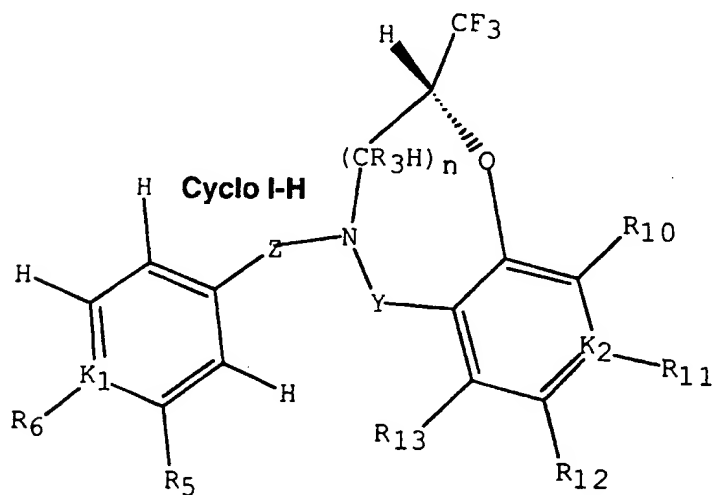


R_2 is hydrido;

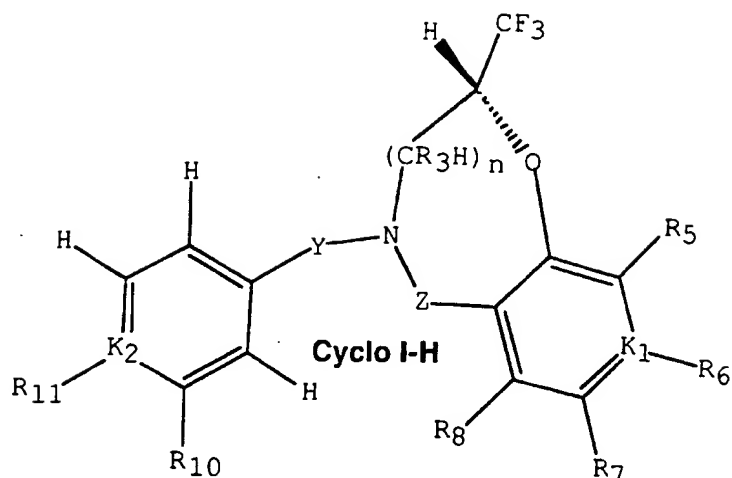
R_2 can be phenyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

- 5 R_3 is selected from the group consisting of hydrido, methyl, trifluoromethyl, and difluoromethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 .

10 In another embodiment of compounds of Formulas I-H or I-C, the compounds correspond to the Cyclo I-H Formulas:



and

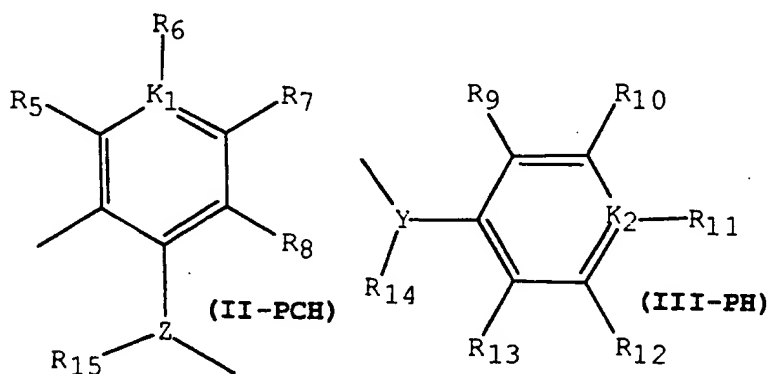


wherein:

K_1 and K_2 are independently selected from the group consisting of C
5 and N;

n is an integer selected from 1 through 3;

R_1 is selected from the group consisting of haloalkyl, haloalkenyl,
haloalkoxymethyl, and haloalkenyloxymethyl with the proviso that R_1 has a
higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and
10 $(CHR_3)_n-N(Apch)Qph$ wherein Apch is Formula (II-PCH) and Qph is
Formula (III-PH);



R_2 is hydrido;

R_2 is selected from the group consisting of aryl, aralkyl, alkyl, alkenyl, alkoxyalkyl, alkenyloxyalkyl, haloalkyl, haloalkenyl, halocycloalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, heteroarylalkyl, dicyanoalkyl, and carboalkoxycyanoalkyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Apch)Qph$;

R_3 is selected from the group consisting of hydrido, hydroxy, halo, cyano, aryl, aralkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, heteroaryl, alkenyloxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, carboxamide, and carboxamidoalkyl with the provisos that $(CHR_3)_n-N(Apch)Qph$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than

R_2 ;

Y is selected from the group consisting of a covalent single bond and $(C(R_{14})_2)_q$ wherein q is an integer selected from 1 through 2;

R_{14} is selected from the group consisting of hydrido, hydroxy, cyano, hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkynyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, carboalkoxy, carboxamide, and carboxamidoalkyl;

Z is selected from the group consisting of covalent single bond, $(C(R_{15})_2)_q$ wherein q is an integer selected from 1 through 2, and

$(CH(R_{15}))_j-W-(CH(R_{15}))_k$ wherein j and k are integers independently selected from 0 through 1;

W is selected from the group consisting of O, C(O), S, S(O), and $S(O)_2$;

- R₁₅ is selected from the group consisting of hydrido, cyano, hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkynyl, alkoxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl, carboalkoxycyanoalkyl, carboalkoxy, carboxamide, and carboxamidoalkyl;

R₈, R₉, and R₁₃ are independently selected from the group consisting of hydrido, halo, haloalkyl, and alkyl ;

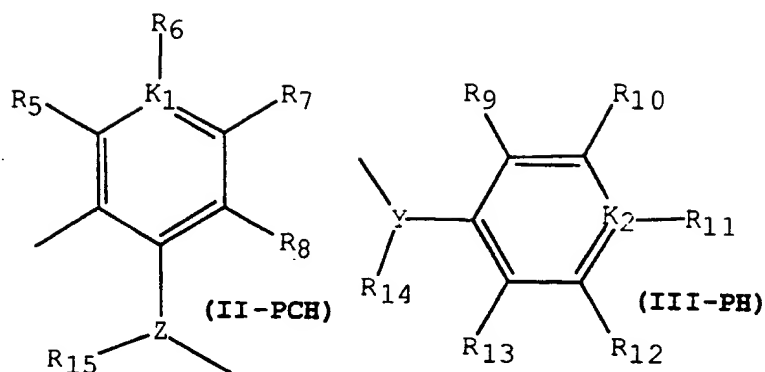
- R₅, R₆, R₇, R₁₀, R₁₁, and R₁₂ are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroaryl amino, N-heteroaryl amino-N-alkyl amino, heteroaryl aminoalkyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxylalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro, lower alkyl amino, alkylthio, alkylthioalkyl, aryl amino, aralkyl amino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, heteroaryl sulfinylalkyl, heteroaryl sulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, alkylsulfonylamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonylamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroaryl sulfinyl, heteroaryl sulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl,

partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxy-carboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido,
 5 carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl;

R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} can be independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3
 10 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no
 15 more than one of the group consisting of spacer pairs R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , can be used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} can be used at the same time.

20 In an embodiment of compounds of Formula Cyclo I-H, n is the integer 1;

R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a
 25 higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Apch)Qph$ wherein $Apch$ is Formula (II-PCH) and Qph is Formula (III-PH);



R₂ is hydrido;

R₂ is selected from the group consisting of phenyl,

- 5 4-trifluoromethylphenyl, vinyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R₂ has a lower Cahn-Ingold-Prelog system ranking than both R₁ and (CHR₃)_n-N(Apch)Qph;

R₃ is selected from the group consisting of hydrido, methyl, ethyl,

- 10 vinyl, phenyl, 4-trifluoromethylphenyl, methoxymethyl, trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the provisos that (CHR₃)_n-N(Apch)Qph has a lower Cahn-Ingold-Prelog stereochemical system ranking than R₁ and a higher Cahn-Ingold-Prelog stereochemical system ranking than R₂.

15

In another embodiment of compounds of Formula Cyclo I-H, n is the integer 1;

- R₁ is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, 20 chlorodifluoromethyl, and pentafluoroethyl;

R₂ is hydrido;

- R_3 is selected from the group consisting of hydrido, methyl, ethyl, vinyl, phenyl, 4-trifluoromethylphenyl, methoxymethyl, trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the provisos that $(CHR_3)_n-N(Apch)Qph$ has a lower
- 5 Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 .

In a preferred embodiment of compounds of Formulas I-H, I-C, I-CP, and Cyclo I-H,

- 10 Y is selected from the group consisting of methylene, ethylene, and ethylidene;
 Z is covalent single bond;
- R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro with the proviso that there is no R_4 , R_8 , R_9 , or R_{13} when the embodiment is a compound of Formula Cyclo I-H;
- 15 R_5 and R_{10} are independently selected from the group consisting of 4-aminophenoxy, benzoyl, benzyl, benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-bromo-2-nitrophenoxy, 3-bromobenzyloxy, 4-bromobenzyloxy, 4-bromophenoxy, 5-bromopyrid-2-yloxy,
- 20 4-butoxyphenoxy, chloro, 3-chlorobenzyl, 2-chlorophenoxy, 4-chlorophenoxy, 4-chloro-3-ethylphenoxy, 3-chloro-4-fluorobenzyl, 3-chloro-4-fluorophenyl, 3-chloro-2-fluorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy,
- 25 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy, 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy, 2-cyanopyrid-3-yloxy, 4-cyanophenoxy, cyclobutoxy, cyclobutyl, cyclohexoxy, cyclohexylmethoxy, cyclopentoxo, cyclopentyl, cyclopentylcarbonyl, cyclopropyl, cyclopropylmethoxy, cyclopropoxy,
- 30 2,3-dichlorophenoxy, 2,4-dichlorophenoxy, 2,4-dichlorophenyl,

- 3,5-dichlorophenyl, 3,5-dichlorobenzyl, 3,4-dichlorophenoxy,
 3,4-difluorophenoxy, 2,3-difluorobenzyl, 2,4-difluorobenzyl, 2,4-difluorobenzyl,
 3,4-difluorobenzyl, 2,5-difluorobenzyl, 3,5-difluorophenoxy,
 3,4-difluorophenyl, 3,5-difluorobenzyl, 4-difluoromethoxybenzyl,
 5 2,3-difluorophenoxy, 2,4-difluorophenoxy, 2,5-difluorophenoxy,
 3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy, 3,5-dimethylphenoxy,
 3,4-dimethylphenoxy, 3,4-dimethylbenzyl, 3,4-dimethylbenzyl,
 3,5-dimethylbenzyl, 2,2-dimethylpropoxy, 1,3-dioxan-2-yl,
 1,4-dioxan-2-yl, 1,3-dioxolan-2-yl, ethoxy, 4-ethoxyphenoxy,
 10 4-ethylbenzyl, 3-ethylphenoxy, 4-ethylaminophenoxy,
 3-ethyl-5-methylphenoxy, fluoro, 4-fluoro-3-methylbenzyl,
 4-fluoro-3-methylphenyl, 4-fluoro-3-methylbenzyl, 4-fluorobenzyl,
 2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy,
 3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzyl,
 15 3-fluoro-5-trifluoromethylbenzyl, 4-fluoro-2-trifluoromethylbenzyl,
 4-fluoro-3-trifluoromethylbenzyl, 2-fluorophenoxy, 4-fluorophenoxy,
 2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzyl,
 4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy,
 4-fluoropyrid-2-yl, 2-furyl, 3-furyl, heptafluoropropyl,
 20 1,1,1,3,3,3-hexafluoropropyl, 2-hydroxy-3,3,3-trifluoropropoxy,
 3-iodobenzyl, isobutyl, isobutylamino, isobutoxy, 3-isoxazolyl,
 4-isoxazolyl, 5-isoxazolyl, isopropoxy, isopropyl, 4-isopropylbenzyl,
 3-isopropylphenoxy, 4-isopropylphenoxy, isopropylthio,
 4-isopropyl-3-methylphenoxy, 3-isothiazolyl, 4-isothiazolyl,
 25 5-isothiazolyl, 3-methoxybenzyl, 4-methoxycarbonylbutoxy,
 3-methoxycarbonylprop-2-enyl, 4-methoxyphenyl,
 3-methoxyphenylamino, 4-methoxyphenylamino, 3-methylbenzyl,
 4-methylbenzyl, 3-methylphenoxy, 3-methyl-4-methylthiophenoxy,
 4-methylphenoxy, 1-methylpropoxy, 2-methylpyrid-5-yl,
 30 4-methylthiophenoxy, 2-naphthyl, 2-nitrophenoxy, 4-nitrophenoxy,
 3-nitrophenyl, 4-nitrophenylthio, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl,
 pentafluoroethyl, pentafluoroethylthio, 2,2,3,3,3-pentafluoropropyl,
 1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl, phenoxy,
 phenylamino, 1-phenylethoxy, phenylsulfonyl, 4-propanoylphenoxy,
 35 propoxy, 4-propylphenoxy, 4-propoxyphenoxy, thiophen-3-yl, *sec*-butyl,
 4-*sec*-butylphenoxy, *tert*-butoxy, 3-*tert*-butylphenoxy, 4-*tert*-butylphenoxy,
 1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,

- 2-(5,6,7,8-tetrahydronaphthoxy), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl, thiophen-2-yl, 2,3,5-trifluorobenzyloxy, 2,2,2-trifluoroethoxy, 2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy, 3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 4-trifluoromethoxyphenoxy, trifluoromethyl, 3-trifluoromethylbenzyloxy, 4-trifluoromethylbenzyloxy, 2,4-bis-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl, 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy, 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy, 3-trifluoromethylphenyl, 3-trifluoromethylthiobenzyloxy, 4-trifluoromethylthiobenzyloxy, 2,3,4-trifluorophenoxy, 2,3,4-trifluorophenyl, 2,3,5-trifluorophenoxy, 3,4,5-trimethylphenoxy, 3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, and trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy, trifluoromethyl, and trifluoromethoxy;

- R_7 and R_{12} are independently selected from the group consisting of hydrido, fluoro, and trifluoromethyl.

- In an even more preferred embodiment of compounds of Formulas I-H, I-C, I-CP, and Cyclo I-H,

Y is methylene;

Z is covalent single bond;

- R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro with the proviso that there is no R_4 , R_8 , R_9 , or R_{13} when the embodiment is a compound of Formula Cyclo I-H;

R_5 and R_{10} are independently selected from the group consisting of benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy,

- 3-bromobenzoyloxy, 4-bromophenoxy, 4-butoxyphenoxy,
3-chlorobenzoyloxy, 2-chlorophenoxy, 4-chloro-3-ethylphenoxy,
4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy,
5 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy,
3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy,
4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy,
cyclobutoxy, cyclobutyl, cyclohexylmethoxy, cyclopentoxo, cyclopentyl,
cyclopentylcarbonyl, cyclopropylmethoxy, 2,3-dichlorophenoxy,
10 2,4-dichlorophenoxy, 2,4-dichlorophenyl, 3,5-dichlorophenyl,
3,5-dichlorobenzyl, 3,4-dichlorophenoxy, 3,4-difluorophenoxy,
2,3-difluorobenzoyloxy, 3,5-difluorobenzoyloxy, difluoromethoxy,
3,5-difluorophenoxy, 3,4-difluorophenyl, 2,3-difluorophenoxy,
2,4-difluorophenoxy, 2,5-difluorophenoxy, 3,5-dimethoxyphenoxy,
15 3-dimethylaminophenoxy, 3,4-dimethylbenzoyloxy, 3,5-dimethylbenzoyloxy,
3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 1,3-dioxolan-2-yl,
3-ethylbenzoyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy,
3-ethyl-5-methylphenoxy, 4-fluoro-3-methylbenzyl, 4-fluorobenzoyloxy,
2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy,
20 3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzoyloxy,
3-fluoro-5-trifluoromethylbenzoyloxy, 2-fluorophenoxy, 4-fluorophenoxy,
2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzoyloxy,
4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy, 2-furyl, 3-furyl,
heptafluoropropyl, 1,1,1,3,3,3-hexafluoropropyl,
25 2-hydroxy-3,3,3-trifluoropropoxy, isobutoxy, isobutyl, 3-isoxazolyl,
4-isoxazolyl, 5-isoxazolyl, isopropoxy,
3-isopropylbenzoyloxy, 3-isopropylphenoxy, isopropylthio,
4-isopropyl-3-methylphenoxy, 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl,
3-methoxybenzyl, 4-methoxyphenylamino, 3-methylbenzoyloxy,
30 4-methylbenzoyloxy, 3-methylphenoxy, 3-methyl-4-methylthiophenoxy,
4-methylphenoxy, 1-methylpropoxy, 2-methylpyrid-5-yloxy,
4-methylthiophenoxy, 2-naphthyl, 2-nitrophenoxy, 4-nitrophenoxy,
3-nitrophenyl, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl, pentafluoroethyl,
pentafluoroethylthio, 2,2,3,3,3-pentafluoropropyl,
35 1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl, phenoxy,
phenylamino, 1-phenylethoxy, 4-propylphenoxy, 4-propoxyphenoxy,

- thiophen-3-yl, tert -butoxy, 3-tert -butylphenoxy, 4-tert -butylphenoxy,
 1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
 2-(5,6,7,8-tetrahydronaphthylthio), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl,
 5 thiophen-2-yl, 2,2,2-trifluoroethoxy, 2,2,2-trifluoroethyl,
 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
 3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
 4-trifluoromethoxyphenoxy, 3-trifluoromethoxyphenoxy, trifluoromethyl,
 3-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,
 10 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
 3-trifluoromethylphenyl, 2,3,4-trifluorophenoxy, 2,3,5-trifluorophenoxy,
 3,4,5-trimethylphenoxy, 3-difluoromethoxyphenoxy,
 3-pentafluoroethylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy,
 15 3-trifluoromethylthiophenoxy, 3-trifluoromethylthiobenzyloxy, and
 trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of
 chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy, and
 trifluoromethyl;

- 20 R_7 and R_{12} are independently selected from the group consisting of
 hydrido, fluoro, and trifluoromethyl.

- In a most preferred embodiment of compounds of Formulas I-H, I-C,
 25 I-CP, and Cyclo I-H,

Y is methylene;

Z is covalent single bond;

- R_4 , R_8 , R_9 , and R_{13} are independently selected from the group
 consisting of hydrido and fluoro with the proviso that there is no R_4 , R_8 , R_9 ,
 30 or R_{13} when the embodiment is a compound of Formula Cyclo I-H;

R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy,
 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-

difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3-ethylphenoxy, 3-ethyl-5-methylphenoxy,
 4-fluoro-3-methylphenoxy, 4-fluorophenoxy, 3-isopropylphenoxy,
 3-methylphenoxy, 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy,
 5 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthylthio),
 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy,
 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R₁₀ is selected from the group consisting of cyclopentyl, 1,1,2,2-tetrafluoroethoxy, 2-furyl, 1,1-bis-trifluoromethyl-1-hydroxymethyl, isobutyl,
 10 isopropoxy, pentafluoroethyl, trifluoromethoxy, trifluoromethyl, and trifluoromethylthio;

R₆ and R₁₁ are independently selected from the group consisting of fluoro and hydrido;

R₇ and R₁₂ are independently selected from the group consisting of
 15 hydrido and fluoro.

DEFINITIONS

The use of generic terms in the description of the compounds are herein defined for clarity.

Standard single letter elemental symbols are used to represent specific types
 20 of atoms unless otherwise defined. The symbol "C" represents a carbon atom. The symbol "O" represents an oxygen atom. The symbol "N" represents a nitrogen atom. The symbol "P" represents a phosphorus atom. The symbol "S" represents a sulfur atom. The symbol "H" represents a hydrogen atom. Double letter elemental symbols are used as defined for the elements of the periodical table
 25 (i.e., Cl represents chlorine, Se represents selenium, etc.).

As utilized herein, the term "alkyl", either alone or within other terms such as "haloalkyl" and "alkylthio", means an acyclic alkyl radical containing from 1 to about 10, preferably from 1 to about 8 carbon atoms and more preferably 1 to about 6 carbon atoms. Said alkyl radicals may be optionally substituted with
 30 groups as defined below. Examples of such radicals include methyl, ethyl, chloroethyl, hydroxyethyl, n-propyl, oxopropyl, isopropyl, n-butyl, cyanobutyl, isobutyl, sec-butyl, tert-butyl, pentyl, aminopentyl, iso-amyl, hexyl, octyl and the like.

The term "alkenyl" refers to an unsaturated, acyclic hydrocarbon radical in so much as it contains at least one double bond. Such alkenyl radicals contain from about 2 to about 10 carbon atoms, preferably from about 2 to about 8 carbon atoms and more preferably 2 to about 6 carbon atoms. Said alkenyl radicals may be optionally substituted with groups as defined below. Examples of suitable alkenyl radicals include propenyl, 2-chloropropenyl, buten-1-yl, isobutenyl, penten-1-yl, 2-2-methylbuten-1-yl, 3-methylbuten-1-yl, hexen-1-yl, 3-hydroxyhexen-1-yl, hepten-1-yl, and octen-1-yl, and the like.

The term "alkynyl" refers to an unsaturated, acyclic hydrocarbon radical in so much as it contains one or more triple bonds, such radicals containing about 2 to about 10 carbon atoms, preferably having from about 2 to about 8 carbon atoms and more preferably having 2 to about 6 carbon atoms. Said alkynyl radicals may be optionally substituted with groups as defined below. Examples of suitable alkynyl radicals include ethynyl, propynyl, hydroxypropynyl, butyn-1-yl, butyn-2-yl, pentyn-1-yl, pentyn-2-yl, 4-methoxypentyn-2-yl, 3-methylbutyn-1-yl, hexyn-1-yl, hexyn-2-yl, hexyn-3-yl, 3,3-dimethylbutyn-1-yl radicals and the like.

The term "hydrido" denotes a single hydrogen atom (H). This hydrido radical may be attached, for example, to an oxygen atom to form a "hydroxyl" radical, one hydrido radical may be attached to a carbon atom to form a "methine" radical ($=CH-$), or two hydrido radicals may be attached to a carbon atom to form a "methylene" ($-CH_2-$) radical.

The term "carbon" radical denotes a carbon atom without any covalent bonds and capable of forming four covalent bonds.

The term "cyano" radical denotes a carbon radical having three of four covalent bonds shared by a nitrogen atom.

The term "hydroxyalkyl" embraces radicals wherein any one or more of the alkyl carbon atoms is substituted with a hydroxyl as defined above. Specifically embraced are monohydroxyalkyl, dihydroxyalkyl and polyhydroxyalkyl radicals.

The term "alkanoyl" embraces radicals wherein one or more of the terminal alkyl carbon atoms are substituted with one or more carbonyl radicals as defined below. Specifically embraced are monocarbonylalkyl and dicarbonylalkyl radicals. Examples of monocarbonylalkyl radicals include formyl, acetyl, and pentanoyl. Examples of dicarbonylalkyl radicals include oxalyl, malonyl, and succinyl.

The term "alkylene" radical denotes linear or branched radicals having from 1 to about 10 carbon atoms and having attachment points for two or more covalent bonds. Examples of such radicals are methylene, ethylene, ethylidene, methylethylene, and isopropylidene.

5 The term "alkenylene" radical denotes linear or branched radicals having from 2 to about 10 carbon atoms, at least one double bond, and having attachment points for two or more covalent bonds. Examples of such radicals are 1,1-vinylidene ($\text{CH}_2=\text{C}$), 1,2-vinylidene ($-\text{CH}=\text{CH}-$), and 1,4-butadienyl ($-\text{CH}=\text{CH}-\text{CH}=\text{CH}-$).

10 The term "halo" means halogens such as fluorine, chlorine, bromine or iodine atoms.

 The term "haloalkyl" embraces radicals wherein any one or more of the alkyl carbon atoms is substituted with halo as defined above. Specifically embraced are monohaloalkyl, dihaloalkyl and polyhaloalkyl radicals. A
15 monohaloalkyl radical, for one example, may have either a bromo, chloro or a fluoro atom within the radical. Dihalo radicals may have two or more of the same halo atoms or a combination of different halo radicals and polyhaloalkyl radicals may have more than two of the same halo atoms or a combination of different halo radicals. More preferred haloalkyl radicals are "lower haloalkyl"
20 radicals having one to about six carbon atoms. Examples of such haloalkyl radicals include fluoromethyl, difluoromethyl, trifluoromethyl, chloromethyl, dichloromethyl, trichloromethyl, trifluoroethyl, pentafluoroethyl, heptafluoropropyl, difluorochloromethyl, dichlorofluoromethyl, difluoroethyl, difluoropropyl, dichloroethyl and dichloropropyl.

25 The term "hydroxyhaloalkyl" embraces radicals wherein any one or more of the haloalkyl carbon atoms is substituted with hydroxy as defined above. Examples of "hydroxyhaloalkyl" radicals include hexafluorohydroxypropyl.

 The term "haloalkylene radical" denotes alkylene radicals wherein any
30 one or more of the alkylene carbon atoms is substituted with halo as defined above. Dihalo alkylene radicals may have two or more of the same halo atoms or a combination of different halo radicals and polyhaloalkylene radicals may have more than two of the same halo atoms or a combination of different halo radicals. More preferred haloalkylene radicals are "lower haloalkylene"
35 radicals having one to about six carbon atoms. Examples of "haloalkylene" radicals include difluoromethylene, tetrafluoroethylene, tetrachloroethylene,

alkyl substituted monofluoromethylene, and aryl substituted trifluoromethylene.

5 The term "haloalkenyl" denotes linear or branched radicals having from 1 to about 10 carbon atoms and having one or more double bonds wherein any one or more of the alkenyl carbon atoms is substituted with halo as defined above. Dihaloalkenyl radicals may have two or more of the same halo atoms or a combination of different halo radicals and polyhaloalkenyl radicals may have more than two of the same halo atoms or a combination of different halo radicals.

10 The terms "alkoxy" and "alkoxyalkyl" embrace linear or branched oxy-containing radicals each having alkyl portions of one to about ten carbon atoms, such as methoxy radical. The term "alkoxyalkyl" also embraces alkyl radicals having one or more alkoxy radicals attached to the alkyl radical, that is, to form monoalkoxyalkyl and dialkoxyalkyl radicals. More preferred alkoxy
15 radicals are "lower alkoxy" radicals having one to six carbon atoms. Examples of such radicals include methoxy, ethoxy, propoxy, butoxy, isopropoxy and *tert*-butoxy alkyls. The "alkoxy" radicals may be further substituted with one or more halo atoms, such as fluoro, chloro or bromo, to provide "haloalkoxy" and "haloalkoxyalkyl" radicals. Examples of such haloalkoxy radicals include
20 fluoromethoxy, chloromethoxy, trifluoromethoxy, difluoromethoxy, trifluoroethoxy, fluoroethoxy, tetrafluoroethoxy, pentafluoroethoxy, and fluoropropoxy. Examples of such haloalkoxyalkyl radicals include fluoromethoxymethyl, chloromethoxyethyl, trifluoromethoxymethyl, difluoromethoxyethyl, and trifluoroethoxymethyl.

25 The terms "alkenyloxy" and "alkenyloxyalkyl" embrace linear or branched oxy-containing radicals each having alkenyl portions of two to about ten carbon atoms, such as ethenyloxy or propenyloxy radical. The term "alkenyloxyalkyl" also embraces alkenyl radicals having one or more alkenyloxy radicals attached to the alkyl radical, that is, to form
30 monoalkenyloxyalkyl and dialkenyloxyalkyl radicals. More preferred alkenyloxy radicals are "lower alkenyloxy" radicals having two to six carbon atoms. Examples of such radicals include ethenyloxy, propenyloxy, butenyloxy, and isopropenyloxy alkyls. The "alkenyloxy" radicals may be further substituted with one or more halo atoms, such as fluoro, chloro or
35 bromo, to provide "haloalkenyloxy" radicals. Examples of such radicals include trifluoroethenyloxy, fluoroethenyloxy, difluoroethenyloxy, and fluoropropenyloxy.

The term "haloalkoxyalkyl" also embraces alkyl radicals having one or more haloalkoxy radicals attached to the alkyl radical, that is, to form monohaloalkoxyalkyl and dihaloalkoxyalkyl radicals. The term "haloalkenyloxy" also embraces oxygen radicals having one or more haloalkenyloxy radicals attached to the oxygen radical, that is, to form monohaloalkenyloxy and dihaloalkenyloxy radicals. The term "haloalkenyloxyalkyl" also embraces alkyl radicals having one or more haloalkenyloxy radicals attached to the alkyl radical, that is, to form monohaloalkenyloxyalkyl and dihaloalkenyloxyalkyl radicals.

10 The term "alkylenedioxy" radicals denotes alkylene radicals having at least two oxygens bonded to a single alkylene group. Examples of "alkylenedioxy" radicals include methylenedioxy, ethylenedioxy, alkylsubstituted methylenedioxy, and arylsubstituted methylenedioxy. The term "haloalkylenedioxy" radicals denotes haloalkylene radicals having at least two
15 oxy groups bonded to a single haloalkyl group. Examples of "haloalkylenedioxy" radicals include difluoromethylenedioxy, tetrafluoroethylenedioxy, tetrachloroethylenedioxy, alkylsubstituted monofluoromethylenedioxy, and arylsubstituted monofluoromethylenedioxy.

The term "aryl", alone or in combination, means a carbocyclic aromatic
20 system containing one, two or three rings wherein such rings may be attached together in a pendant manner or may be fused. The term "fused" means that a second ring is present (ie, attached or formed) by having two adjacent atoms in common (ie, shared) with the first ring. The term "fused" is equivalent to the term "condensed". The term "aryl" embraces aromatic radicals such as phenyl,
25 naphthyl, tetrahydronaphthyl, indane and biphenyl.

The term "perhaloaryl" embraces aromatic radicals such as phenyl, naphthyl, tetrahydronaphthyl, indane and biphenyl wherein the aryl radical is substituted with 3 or more halo radicals as defined below.

The term "heterocyclyl" embraces saturated, partially saturated and
30 unsaturated heteroatom-containing ring-shaped radicals having from 5 through 15 ring members selected from carbon, nitrogen, sulfur and oxygen, wherein at least one ring atom is a heteroatom. Heterocyclyl radicals may contain one, two or three rings wherein such rings may be attached in a pendant manner or may be fused. Examples of saturated heterocyclic radicals include saturated 3
35 to 6-membered heteromonocyclic group containing 1 to 4 nitrogen atoms [e.g. pyrrolidinyl, imidazolidinyl, piperidino, piperazinyl, etc.]; saturated 3 to 6-membered heteromonocyclic group containing 1 to 2 oxygen atoms and 1 to 3

nitrogen atoms [e.g. morpholinyl, etc.]; saturated 3 to 6-membered heteromonocyclic group containing 1 to 2 sulfur atoms and 1 to 3 nitrogen atoms [e.g., thiazolidinyl, etc.]. Examples of partially saturated heterocyclyl radicals include dihydrothiophene, dihydropyran, dihydrofuran and dihydrothiazole. Examples of unsaturated heterocyclic radicals, also termed "heteroaryl" radicals, include unsaturated 5 to 6 membered heteromonocyclyl group containing 1 to 4 nitrogen atoms, for example, pyrrolyl, pyrrolinyl, imidazolyl, pyrazolyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazolyl [e.g., 4H-1,2,4-triazolyl, 1H-1,2,3-triazolyl, 2H-1,2,3-triazolyl, etc.] tetrazolyl [e.g. 1H-tetrazolyl, 2H-tetrazolyl, etc.], etc.; unsaturated condensed heterocyclic group containing 1 to 5 nitrogen atoms, for example, indolyl, isoindolyl, indolizinyl, benzimidazolyl, quinolyl, isoquinolyl, indazolyl, benzotriazolyl, tetrazolopyridazinyl [e.g., tetrazolo [1,5-b]pyridazinyl, etc.], etc.; unsaturated 3 to 6-membered heteromonocyclic group containing an oxygen atom, for example, pyranyl, 2-furyl, 3-furyl, etc.; unsaturated 5 to 6-membered heteromonocyclic group containing a sulfur atom, for example, 2-thienyl, 3-thienyl, etc.; unsaturated 5- to 6-membered heteromonocyclic group containing 1 to 2 oxygen atoms and 1 to 3 nitrogen atoms, for example, oxazolyl, isoxazolyl, oxadiazolyl [e.g., 1,2,4-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,5-oxadiazolyl, etc.] etc.; unsaturated condensed heterocyclic group containing 1 to 2 oxygen atoms and 1 to 3 nitrogen atoms [e.g. benzoxazolyl, benzoxadiazolyl, etc.]; unsaturated 5 to 6-membered heteromonocyclic group containing 1 to 2 sulfur atoms and 1 to 3 nitrogen atoms, for example, thiazolyl, thiadiazolyl [e.g., 1,2,4-thiadiazolyl, 1,3,4-thiadiazolyl, 1,2,5-thiadiazolyl, etc.] etc.; unsaturated condensed heterocyclic group containing 1 to 2 sulfur atoms and 1 to 3 nitrogen atoms [e.g., benzothiazolyl, benzothiadiazolyl, etc.] and the like. The term also embraces radicals where heterocyclic radicals are fused with aryl radicals. Examples of such fused bicyclic radicals include benzofuran, benzothiophene, and the like.

Said "heterocyclyl" group may have 1 to 3 substituents as defined below. Preferred heterocyclic radicals include five to twelve membered fused or unfused radicals. Non-limiting examples of heterocyclic radicals include pyrrolyl, pyridinyl, pyridyloxy, pyrazolyl, triazolyl, pyrimidinyl, pyridazinyl, oxazolyl, thiazolyl, imidazolyl, indolyl, thiophenyl, furanyl, tetrazolyl, 2-pyrrolinyl, 3-pyrrolinyl, pyrrolindinyl, 1,3-dioxolanyl, 2-imidazoliny, imidazolidinyl, 2-pyrazolinyl, pyrazolidinyl, isoxazolyl, isothiazolyl, 1,2,3-oxadiazolyl, 1,2,3-triazolyl, 1,3,4-thiadiazolyl, 2H-pyranyl, 4H-pyranyl,

piperidinyl, 1,4-dioxanyl, morpholinyl, 1,4-dithianyl, thiomorpholinyl, pyrazinyl, piperazinyl, 1,3,5-triazinyl, 1,3,5-trithianyl, benzo(b)thiophenyl, benzimidazolyl, quinolinyl, tetraazolyl, and the like.

The term "sulfonyl", whether used alone or linked to other terms such as alkylsulfonyl, denotes respectively divalent radicals $-SO_2-$. "Alkylsulfonyl",
embraces alkyl radicals attached to a sulfonyl radical, where alkyl is defined as above. "Alkylsulfonylalkyl", embraces alkylsulfonyl radicals attached to an alkyl radical, where alkyl is defined as above. "Haloalkylsulfonyl", embraces haloalkyl radicals attached to a sulfonyl radical, where haloalkyl is defined as above. "Haloalkylsulfonylalkyl", embraces haloalkylsulfonyl radicals attached to an alkyl radical, where alkyl is defined as above. The term "aminosulfonyl" denotes an amino radical attached to a sulfonyl radical.

The term "sulfinyl", whether used alone or linked to other terms such as alkylsulfinyl, denotes respectively divalent radicals $-S(O)-$. "Alkylsulfinyl",
embraces alkyl radicals attached to a sulfinyl radical, where alkyl is defined as above. "Alkylsulfinylalkyl", embraces alkylsulfinyl radicals attached to an alkyl radical, where alkyl is defined as above. "Haloalkylsulfinyl", embraces haloalkyl radicals attached to a sulfinyl radical, where haloalkyl is defined as above. "Haloalkylsulfinylalkyl", embraces haloalkylsulfinyl radicals attached to an alkyl radical, where alkyl is defined as above.

The term "aralkyl" embraces aryl-substituted alkyl radicals. Preferable aralkyl radicals are "lower aralkyl" radicals having aryl radicals attached to alkyl radicals having one to six carbon atoms. Examples of such radicals include benzyl, diphenylmethyl, triphenylmethyl, phenylethyl and diphenylethyl. The terms benzyl and phenylmethyl are interchangeable.

The term "heteroaralkyl" embraces heteroaryl-substituted alkyl radicals wherein the heteroaralkyl radical may be additionally substituted with three or more substituents as defined above for aralkyl radicals. The term "perhaloaralkyl" embraces aryl-substituted alkyl radicals wherein the aralkyl radical is substituted with three or more halo radicals as defined above.

The term "aralkylsulfinyl", embraces aralkyl radicals attached to a sulfinyl radical, where aralkyl is defined as above. "Aralkylsulfinylalkyl", embraces aralkylsulfinyl radicals attached to an alkyl radical, where alkyl is defined as above.

The term "aralkylsulfonyl", embraces aralkyl radicals attached to a sulfonyl radical, where aralkyl is defined as above. "Aralkylsulfonylalkyl",

embraces aralkylsulfonyl radicals attached to an alkyl radical, where alkyl is defined as above.

The term "cycloalkyl" embraces radicals having three to ten carbon atoms. More preferred cycloalkyl radicals are "lower cycloalkyl" radicals having three to seven carbon atoms. Examples include radicals such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl. The term "cycloalkylalkyl" embraces cycloalkyl-substituted alkyl radicals. Preferable cycloalkylalkyl radicals are "lower cycloalkylalkyl" radicals having cycloalkyl radicals attached to alkyl radicals having one to six carbon atoms. Examples of such radicals include cyclohexylhexyl. The term "cycloalkenyl" embraces radicals having three to ten carbon atoms and one or more carbon-carbon double bonds. Preferred cycloalkenyl radicals are "lower cycloalkenyl" radicals having three to seven carbon atoms. Examples include radicals such as cyclobutenyl, cyclopentenyl, cyclohexenyl and cycloheptenyl. The term "halocycloalkyl" embraces radicals wherein any one or more of the cycloalkyl carbon atoms is substituted with halo as defined above. Specifically embraced are monohalocycloalkyl, dihalocycloalkyl and polyhalocycloalkyl radicals. A monohalocycloalkyl radical, for one example, may have either a bromo, chloro or a fluoro atom within the radical. Dihalo radicals may have two or more of the same halo atoms or a combination of different halo radicals and polyhalocycloalkyl radicals may have more than two of the same halo atoms or a combination of different halo radicals. More preferred halocycloalkyl radicals are "lower halocycloalkyl" radicals having three to about eight carbon atoms. Examples of such halocycloalkyl radicals include fluorocyclopropyl, difluorocyclobutyl, trifluorocyclopentyl, tetrafluorocyclohexyl, and dichlorocyclopropyl. The term "halocycloalkenyl" embraces radicals wherein any one or more of the cycloalkenyl carbon atoms is substituted with halo as defined above. Specifically embraced are monohalocycloalkenyl, dihalocycloalkenyl and polyhalocycloalkenyl radicals.

The term "cycloalkoxy" embraces cycloalkyl radicals attached to an oxy radical. Examples of such radicals includes cyclohexoxy and cyclopentoxy. The term "cycloalkoxyalkyl" also embraces alkyl radicals having one or more cycloalkoxy radicals attached to the alkyl radical, that is, to form monocycloalkoxyalkyl and dicycloalkoxyalkyl radicals. Examples of such radicals include cyclohexoxyethyl. The "cycloalkoxy" radicals may be further substituted with one or more halo atoms, such as fluoro, chloro or bromo, to provide "halocycloalkoxy" and "halocycloalkoxyalkyl" radicals.

The term "cycloalkylalkoxy" embraces cycloalkyl radicals attached to an alkoxy radical. Examples of such radicals includes cyclohexylmethoxy and cyclopentylmethoxy.

5 The term "cycloalkenyloxy" embraces cycloalkenyl radicals attached to an oxy radical. Examples of such radicals includes cyclohexenyloxy and cyclopentenyloxy. The term "cycloalkenyloxyalkyl" also embraces alkyl radicals having one or more cycloalkenyloxy radicals attached to the alkyl radical, that is, to form monocycloalkenyloxyalkyl and dicyclocycloalkenyloxyalkyl radicals. Examples of such radicals include cyclohexenyloxyethyl. The
10 "cycloalkenyloxy" radicals may be further substituted with one or more halo atoms, such as fluoro, chloro or bromo, to provide "halocycloalkenyloxy" and "halocycloalkenyloxyalkyl" radicals.

The term "cycloalkylenedioxy" radicals denotes cycloalkylene radicals having at least two oxygens bonded to a single cycloalkylene group. Examples
15 of "alkylenedioxy" radicals include 1,2-dioxycyclohexylene.

The term "cycloalkylsulfinyl", embraces cycloalkyl radicals attached to a sulfinyl radical, where cycloalkyl is defined as above. "Cycloalkylsulfinylalkyl", embraces cycloalkylsulfinyl radicals attached to an alkyl radical, where alkyl is defined as above. The term "Cycloalkylsulfonyl",
20 embraces cycloalkyl radicals attached to a sulfonyl radical, where cycloalkyl is defined as above. "Cycloalkylsulfonylalkyl", embraces cycloalkylsulfonyl radicals attached to an alkyl radical, where alkyl is defined as above.

The term "cycloalkylalkanoyl" embraces radicals wherein one or more of the cycloalkyl carbon atoms are substituted with one or more carbonyl
25 radicals as defined below. Specifically embraced are monocarbonylcycloalkyl and dicarbonylcycloalkyl radicals. Examples of monocarbonylcycloalkyl radicals include cyclohexylcarbonyl, cyclohexylacetyl, and cyclopentylcarbonyl. Examples of dicarbonylcycloalkyl radicals include 1,2-dicarbonylcyclohexane..

30 The term "alkylthio" embraces radicals containing a linear or branched alkyl radical, of one to ten carbon atoms, attached to a divalent sulfur atom. More preferred alkylthio radicals are "lower alkylthio" radicals having one to six carbon atoms. An example of "lower alkylthio" is methylthio ($\text{CH}_3\text{-S-}$). The "alkylthio" radicals may be further substituted with one or more halo
35 atoms, such as fluoro, chloro or bromo, to provide "haloalkylthio" radicals. Examples of such radicals include fluoromethylthio, chloromethylthio,

trifluoromethylthio, difluoromethylthio, trifluoroethylthio, fluoroethylthio, tetrafluoroethylthio, pentafluoroethylthio, and fluoropropylthio.

The term "alkyl aryl amino" embraces radicals containing a linear or branched alkyl radical, of one to ten carbon atoms, and one aryl radical both
5 attached to an amino radical. Examples include N-methyl-4-methoxyaniline, N-ethyl-4-methoxyaniline, and N-methyl-4-trifluoromethoxyaniline.

The terms alkylamino denotes "monoalkylamino" and "dialkylamino" containing one or two alkyl radicals, respectively, attached to an amino radical.

The terms arylamino denotes "monoarylamino" and "diarylamino"
10 containing one or two aryl radicals, respectively, attached to an amino radical. Examples of such radicals include N-phenylamino and N-naphthylamino.

The term "aralkylamino", embraces aralkyl radicals attached to an amino radical, where aralkyl is defined as above. The term aralkylamino denotes "monoaralkylamino" and "diaralkylamino" containing one or two
15 aralkyl radicals, respectively, attached to an amino radical. The term aralkylamino further denotes "monoaralkyl monoalkylamino" containing one aralkyl radical and one alkyl radical attached to an amino radical.

The term "arylsulfinyl" embraces radicals containing an aryl radical, as defined above, attached to a divalent S(=O) atom. The term "arylsulfinylalkyl"
20 denotes arylsulfinyl radicals attached to a linear or branched alkyl radical, of one to ten carbon atoms.

The term "arylsulfonyl", embraces aryl radicals attached to a sulfonyl radical, where aryl is defined as above. "arylsulfonylalkyl", embraces arylsulfonyl radicals attached to an alkyl radical, where alkyl is defined as
25 above. The term "heteroarylsulfinyl" embraces radicals containing an heteroaryl radical, as defined above, attached to a divalent S(=O) atom. The term "heteroarylsulfinylalkyl" denotes heteroarylsulfinyl radicals attached to a linear or branched alkyl radical, of one to ten carbon atoms. The term "Heteroarylsulfonyl", embraces heteroaryl radicals attached to a sulfonyl
30 radical, where heteroaryl is defined as above. "Heteroarylsulfonylalkyl", embraces heteroarylsulfonyl radicals attached to an alkyl radical, where alkyl is defined as above.

The term "aryloxy" embraces aryl radicals, as defined above, attached to an oxygen atom. Examples of such radicals include phenoxy, 4-chloro-3-
35 ethylphenoxy, 4-chloro-3-methylphenoxy, 3-chloro-4-ethylphenoxy, 3,4-dichlorophenoxy, 4-methylphenoxy, 3-trifluoromethoxyphenoxy, 3-trifluoromethylphenoxy, 4-fluorophenoxy, 3,4-dimethylphenoxy, 5-bromo-2-

fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-fluoro-3-methylphenoxy, 5,6,7,8-tetrahydronaphthylloxy, 3-isopropylphenoxy, 3-cyclopropylphenoxy, 3-ethylphenoxy, 4-*tert*-butylphenoxy, 3-pentafluoroethylphenoxy, and 3-(1,1,2,2-tetrafluoroethoxy)phenoxy.

5 The term "aroyl" embraces aryl radicals, as defined above, attached to an carbonyl radical as defined above. Examples of such radicals include benzoyl and toluoyl.

10 The term "aralkanoyl" embraces aralkyl radicals, as defined herein, attached to an carbonyl radical as defined above. Examples of such radicals include, for example, phenylacetyl.

15 The term "aralkoxy" embraces oxy-containing aralkyl radicals attached through an oxygen atom to other radicals. More preferred aralkoxy radicals are "lower aralkoxy" radicals having phenyl radicals attached to lower alkoxy radical as described above. Examples of such radicals include benzyloxy, 1-phenylethoxy, 3-trifluoromethoxybenzyloxy, 3-trifluoromethylbenzyloxy, 3,5-difluorobenzyloxy, 3-bromobenzyloxy, 4-propylbenzyloxy, 2-fluoro-3-trifluoromethylbenzyloxy, and 2-phenylethoxy.

20 The term "aryloxyalkyl" embraces aryloxy radicals, as defined above, attached to an alkyl group. Examples of such radicals include phenoxymethyl.

 The term "haloaryloxyalkyl" embraces aryloxyalkyl radicals, as defined above, wherein one to five halo radicals are attached to an aryloxy group.

 The term "heteroaroyl" embraces heteroaryl radicals, as defined above, attached to an carbonyl radical as defined above. Examples of such radicals include furoyl and nicotiny.

25 The term "heteroaralkanoyl" embraces heteroaralkyl radicals, as defined herein, attached to an carbonyl radical as defined above. Examples of such radicals include, for example, pyridylacetyl and furylbutyryl.

30 The term "heteroaralkoxy" embraces oxy-containing heteroaralkyl radicals attached through an oxygen atom to other radicals. More preferred heteroaralkoxy radicals are "lower heteroaralkoxy" radicals having heteroaryl radicals attached to lower alkoxy radical as described above.

 The term "haloheteroaryloxyalkyl" embraces heteroaryloxyalkyl radicals, as defined above, wherein one to four halo radicals are attached to an heteroaryloxy group.

35 The term "heteroarylamino" embraces heterocyclyl radicals, as defined above, attached to an amino group. Examples of such radicals include pyridylamino.

The term "heteroarylaminomethyl" embraces heteroarylaminomethyl radicals, as defined above, attached to an alkyl group. Examples of such radicals include pyridylmethylamino.

5 The term "heteroaryloxy" embraces heterocyclyl radicals, as defined above, attached to an oxy group. Examples of such radicals include 2-thiophenyloxy, 2-pyrimidyloxy, 2-pyridyloxy, 3-pyridyloxy, and 4-pyridyloxy.

10 The term "heteroaryloxyalkyl" embraces heteroaryloxy radicals, as defined above, attached to an alkyl group. Examples of such radicals include 2-pyridyloxymethyl, 3-pyridyloxyethyl, and 4-pyridyloxymethyl.

The term "arylthio" embraces aryl radicals, as defined above, attached to an sulfur atom. Examples of such radicals include phenylthio.

15 The term "arylthioalkyl" embraces arylthio radicals, as defined above, attached to an alkyl group. Examples of such radicals include phenylthiomethyl.

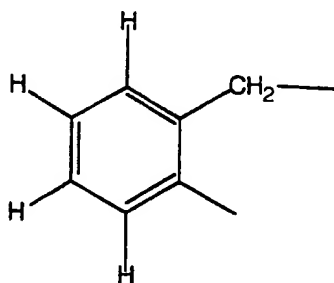
20 The term "alkylthioalkyl" embraces alkylthio radicals, as defined above, attached to an alkyl group. Examples of such radicals include methylthiomethyl. The term "alkoxyalkyl" embraces alkoxy radicals, as defined above, attached to an alkyl group. Examples of such radicals include methoxymethyl.

The term "carbonyl" denotes a carbon radical having two of the four covalent bonds shared with an oxygen atom. The term "carboxy" embraces a hydroxyl radical, as defined above, attached to one of two unshared bonds in a carbonyl group. The term "carboxamide" embraces amino, monoalkylamino, 25 dialkylamino, monocycloalkylamino, alkylcycloalkylamino, and dicycloalkylamino radicals, attached to one of two unshared bonds in a carbonyl group. The term "carboxamidoalkyl" embraces carboxamide radicals, as defined above, attached to an alkyl group. The term "carboxyalkyl" embraces a carboxy radical, as defined above, attached to an alkyl group. The term "carboalkoxy" embraces alkoxy 30 radicals, as defined above, attached to one of two unshared bonds in a carbonyl group. The term "carboaralkoxy" embraces aralkoxy radicals, as defined above, attached to one of two unshared bonds in a carbonyl group. The term "monocarboalkoxyalkyl" embraces one carboalkoxy radical, as defined above, attached to an alkyl group. The term "dicarboalkoxyalkyl" embraces two 35 carboalkoxy radicals, as defined above, attached to an alkylene group. The term "monocyanoalkyl" embraces one cyano radical, as defined above, attached to an alkyl group. The term "dicyanoalkylene" embraces two cyano radicals, as defined

above, attached to an alkyl group. The term "carboalkoxycyanoalkyl" embraces one cyano radical, as defined above, attached to an carboalkoxyalkyl group.

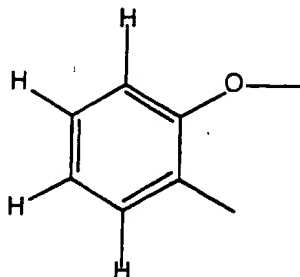
- The term "acyl", alone or in combination, means a carbonyl or thionocarbonyl group bonded to a radical selected from, for example, hydrido, alkyl, alkenyl, alkynyl, haloalkyl, alkoxy, alkoxyalkyl, haloalkoxy, aryl, heterocyclyl, heteroaryl, alkylsulfinylalkyl, alkylsulfonylalkyl, aralkyl, cycloalkyl, cycloalkylalkyl, cycloalkenyl, alkylthio, arylthio, amino, alkylamino, dialkylamino, aralkoxy, arylthio, and alkylthioalkyl. Examples of "acyl" are formyl, acetyl, benzoyl, trifluoroacetyl, phthaloyl, malonyl, nicotinyl, and the like.
- The term "haloalkanoyl" embraces one or more halo radicals, as defined herein, attached to an alkanoyl radical as defined above. Examples of such radicals include, for example, chloroacetyl, trifluoroacetyl, bromopropanoyl, and heptafluorobutanoyl. The term "diacyl", alone or in combination, means having two or more carbonyl or thionocarbonyl groups bonded to a radical selected from, for example, alkylene, alkenylene, alkynylene, haloalkylene, alkoxyalkylene, aryl, heterocyclyl, heteroaryl, aralkyl, cycloalkyl, cycloalkylalkyl, and cycloalkenyl. Examples of "diacyl" are phthaloyl, malonyl, succinyl, adipoyl, and the like.

- The term "benzylidenyl" radical denotes substituted and unsubstituted benzyl groups having attachment points for two covalent bonds. One attachment point is through the methylene of the benzyl group with the other attachment point through an ortho carbon of the phenyl ring. The methylene group is designated for attached to the lowest numbered position. Examples include the base compound benzylidene of structure:



- The term "phenoxyldenyl" radical denotes substituted and unsubstituted phenoxy groups having attachment points for two covalent bonds. One attachment point is through the oxy of the phenoxy group with the other attachment point through an ortho carbon of the phenyl ring. The oxy group is designated for

attached to the lowest numbered position. Examples include the base compound phenoxyldiene of structure:



The term "phosphono" embraces a pentavalent phosphorus attached with two covalent bonds to an oxygen radical. The term "dialkoxyposphono" denotes two alkoxy radicals, as defined above, attached to a phosphono radical with two covalent bonds. The term "diaralkoxyposphono" denotes two aralkoxy radicals, as defined above, attached to a phosphono radical with two covalent bonds. The term "dialkoxyposphonoalkyl" denotes dialkoxyposphono radicals, as defined above, attached to an alkyl radical. The term "diaralkoxyposphonoalkyl" denotes diaralkoxyposphono radicals, as defined above, attached to an alkyl radical.

Said "alkyl", "alkenyl", "alkynyl", "alkanoyl", "alkylene", "alkenylene", "benzylidenyl", "phenoxyldienyl", "hydroxyalkyl", "haloalkyl", "haloalkylene", "haloalkenyl", "alkoxy", "alkenyloxy", "alkenyloxyalkyl", "alkoxyalkyl", "aryl", "perhaloaryl", "haloalkoxy", "haloalkoxyalkyl", "haloalkenyloxy", "haloalkenyloxyalkyl", "alkylenedioxy", "haloalkylenedioxy", "heterocycl", "heteroaryl", "hydroxyhaloalkyl", "alkylsulfonyl", "haloalkylsulfonyl", "alkylsulfonylalkyl", "haloalkylsulfonylalkyl", "alkylsulfinyl", "alkylsulfinylalkyl", "haloalkylsulfinylalkyl", "aralkyl", "heteroaralkyl", "perhaloaralkyl", "aralkylsulfonyl", "aralkylsulfonylalkyl", "aralkylsulfinyl", "aralkylsulfinylalkyl", "cycloalkyl", "cycloalkylalkanoyl", "cycloalkylalkyl", "cycloalkenyl", "halocycloalkyl", "halocycloalkenyl", "cycloalkylsulfinyl", "cycloalkylsulfinylalkyl", "cycloalkylsulfonyl", "cycloalkylsulfonylalkyl", "cycloalkoxy", "cycloalkoxyalkyl", "cycloalkylalkoxy", "cycloalkenyloxy", "cycloalkenyloxyalkyl", "cycloalkylenedioxy", "halocycloalkoxy", "halocycloalkoxyalkyl", "halocycloalkenyloxy", "halocycloalkenyloxyalkyl", "alkylthio", "haloalkylthio", "alkylsulfinyl", "amino", "oxy", "thio", "alkylamino", "arylamino", "aralkylamino", "arylsulfinyl", "arylsulfinylalkyl", "arylsulfonyl", "arylsulfonylalkyl", "heteroarylsulfinyl", "heteroarylsulfinylalkyl".

- “heteroarylsulfonyl”, “heteroarylsulfonylalkyl”, “heteroaryl amino”,
 “heteroaryl aminoalkyl”, “heteroaryloxy”, “heteroaryloxyalkyl”, “aryloxy”,
 “aroyl”, “aralkanoyl”, “aralkoxy”, “aryloxyalkyl”, “haloaryloxyalkyl”,
 “heteroaroyl”, “heteroaralkanoyl”, “heteroaralkoxy”, “heteroaralkoxyalkyl”,
 5 “arylthio”, “arylthioalkyl”, “alkoxyalkyl”, “acyl” and “diacyl” groups defined
 above may optionally have 1 to 5 non-hydrido substituents such as perhaloaralkyl,
 aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl,
 halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl,
 cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroaryl amino, N-heteroaryl amino-
 10 N-alkyl amino, heteroaryl aminoalkyl, heteroaryloxy, heteroaryloxyalkyl,
 haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxyalkyl, heteroaralkoxy,
 cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy,
 cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl,
 halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro,
 15 lower alkyl amino, alkylthio, alkylthioalkyl, aryl amino, aralkyl amino, arylthio,
 arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl,
 arylsulfinylalkyl, arylsulfonylalkyl, heteroarylsulfinylalkyl,
 heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl,
 haloalkylsulfonylalkyl, alkylsulfonylamido, alkylaminosulfonyl, amidosulfonyl,
 20 monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl,
 arylsulfonylamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl,
 arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl,
 alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl,
 alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy,
 25 haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower
 cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy,
 hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl,
 haloalkoxyalkyl, aryl, aralkyl, aryloxy, aralkoxy, aryloxyalkyl, saturated
 heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy,
 30 heteroaryloxyalkyl, arylalkyl, heteroarylalkyl, arylalkenyl, heteroarylalkenyl,
 carboxyalkyl, carboalkoxy, alkoxycarbonyl, carboaralkoxy, carboxamido,
 carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl,
 diaralkoxyphosphono, and diaralkoxyphosphonoalkyl.

The term “spacer” can include a covalent bond and a linear moiety
 35 having a backbone of 1 to 7 continuous atoms. The spacer may have 1 to 7

atoms of a univalent or multi-valent chain. Univalent chains may be constituted by a radical selected from $=C(H)-$, $=C(R_{17})-$, $-O-$, $-S-$, $-S(O)-$,

$-S(O)_2-$, $-NH-$, $-N(R_{17})-$, $-N=$, $-CH(OH)-$, $=C(OH)-$, $-CH(OR_{17})-$,

$=C(OR_{17})-$, and $-C(O)-$ wherein R_{17} is selected from alkyl, alkenyl, alkynyl,

5 aryl, heteroaryl, aralkyl, aryloxyalkyl, alkoxyalkyl, alkylthioalkyl, arylthioalkyl, cycloalkyl, cycloalkylalkyl, haloalkyl, haloalkenyl, haloalkoxyalkyl, perhaloaralkyl, heteroarylalkyl, heteroaryloxyalkyl, heteroarylthioalkyl, and heteroarylalkenyl. Multi-valent chains may consist of

10 a straight chain of 1 or 2 or 3 or 4 or 5 or 6 or 7 atoms or a straight chain of 1 or 2 or 3 or 4 or 5 or 6 atoms with a side chain. The chain may be constituted of one or more radicals selected from: lower alkylene, lower alkenyl, $-O-$, $-O-CH_2-$, $-S-CH_2-$, $-CH_2CH_2-$, ethenyl, $-CH=CH(OH)-$,

$-OCH_2O-$, $-O(CH_2)_2O-$, $-NHCH_2-$, $-OCH(R_{17})O-$, $-O(CH_2CHR_{17})O-$,

$-OCF_2O-$, $-O(CF_2)_2O-$, $-S-$, $-S(O)-$, $-S(O)_2-$, $-N(H)-$, $-N(H)O-$,

15 $-N(R_{17})O-$, $-N(R_{17})-$, $-C(O)-$, $-C(O)NH-$, $-C(O)NR_{17}-$, $-N=$, $-OCH_2-$,

$-SCH_2-$, $S(O)CH_2-$, $-CH_2C(O)-$, $-CH(OH)-$, $=C(OH)-$, $-CH(OR_{17})-$,

$=C(OR_{17})-$, $S(O)_2CH_2-$, and $-NR_{17}CH_2-$ and many other radicals defined

above or generally known or ascertained by one of skill-in-the art. Side chains may include substituents such as 1 to 5 non-hydrido substituents such as

20 perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl,

heteroarylamino, N-heteroarylamino-N-alkylamino, heteroarylaminoalkyl, heteroaryloxy, heteroaryloxyalkyl, haloalkylthio, alkanoyloxy, alkoxy,

25 alkoxyalkyl, haloalkoxyalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl,

cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl,

halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro,

lower alkylamino, alkylthio, alkylthioalkyl, arylamino, aralkylamino, arylthio,

30 arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl,

arylsulfinylalkyl, arylsulfonylalkyl, heteroarylsulfinylalkyl,

heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, 5 diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkenyl, lower cycloalkylalkyl, lower 10 cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, aralkyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, arylalkyl, heteroarylalkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, 15 carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl.

Chiral compounds of the present invention have a hydroxyl group substituent on a chiral carbon of the alkanol and propanol compounds of the 20 present invention specifically in the R-stereoisomeric configuration based on the Cahn-Ingold-Prelog convention for stereoisomeric carbon atoms. The R-stereoisomeric configuration compounds of the present invention may optionally have one or more additional chiral carbons present in each compound. The R-stereoisomeric configuration compounds of the present invention can exist in 25 tautomeric, geometric, and other stereoisomeric forms. The present invention having a hydroxyl group substituent on a chiral carbon of the alkanol and propanol compounds in the R-stereoisomeric configuration contemplates all such forms of said invented compounds, including cis- and trans-geometric isomers, E- and Z-geometric isomers, diastereomers, and other mixtures thereof, as falling within the 30 scope of the invention. Pharmaceutically acceptable salts of such tautomeric, geometric or stereoisomeric forms are also included within the invention. The standard definitions for the Cahn-Ingold-Prelog convention and stereochemical system can be found in Pure Applied Chemistry, 1976, Vol. 45, pages 15-30 and Cahn et al., Angewandte Chemie International Edition English, 1966, Vol. 5, 35 pages 385-415.

The terms "cis" and "trans" denote a form of geometric isomerism in which two carbon atoms connected by a double bond will each have a

hydrogen atom on the same side of the double bond ("cis") or on opposite sides of the double bond ("trans").

Some of the compounds described contain alkenyl groups, and are meant to include both cis and trans or "E" and "Z" geometric forms.

5 Some of the compounds described contain one or more stereocenters in addition to said hydroxyl group substituent on a chiral carbon of the alkanol and propanol compounds in the R-stereoisomeric configuration and are meant to include R, S, and mixtures of R and S forms for each additional stereocenter present.

10 Some of the compounds described herein may contain one or more ketonic or aldehydic carbonyl groups or combinations thereof alone or as part of a heterocyclic ring system. Such carbonyl groups may exist in part or principally in the "keto" form and in part or principally as one or more "enol" forms of each aldehyde and ketone group present. Compounds of the present
15 invention having aldehydic or ketonic carbonyl groups are meant to include both "keto" and "enol" tautomeric forms.

Some of the compounds described herein may contain one or more amide carbonyl groups or combinations thereof alone or as part of a heterocyclic ring system. Such carbonyl groups may exist in part or principally
20 in the "keto" form and in part or principally as one or more "enol" forms of each amide group present. Compounds of the present invention having amidic carbonyl groups are meant to include both "keto" and "enol" tautomeric forms. Said amide carbonyl groups may be both oxo (C=O) and thiono (C=S) in type.

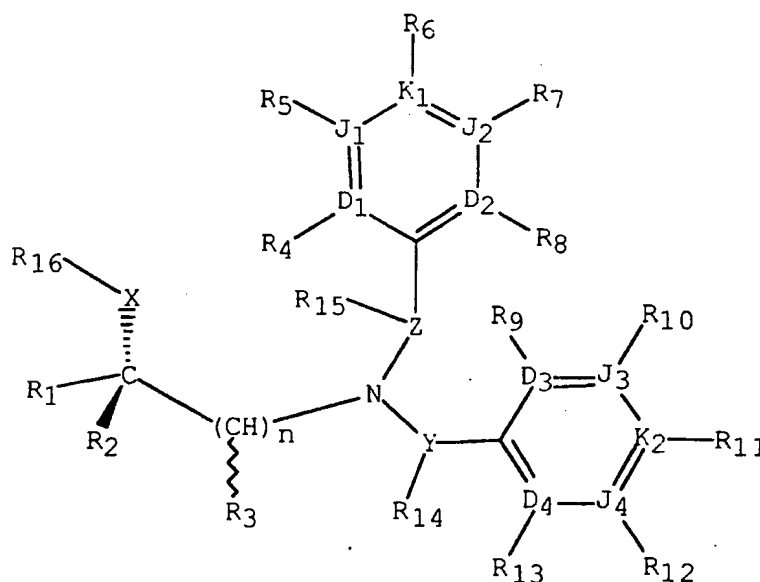
Some of the compounds described herein may contain one or more
25 imine or enamine groups or combinations thereof. Such groups may exist in part or principally in the "imine" form and in part or principally as one or more "enamine" forms of each group present. Compounds of the present invention having said imine or enamine groups are meant to include both "imine" and "enamine" tautomeric forms.

30 The following general synthetic sequences are useful in making the present invention. Abbreviations used in the schemes are as follows: "AA" represents amino acids, "BINAP" represents 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl, "Boc" represents tert-butyloxycarbonyl, "BOP" represents benzotriazol-1-yl-oxy-tris-(dimethylamino), "bu" represents butyl, "dba"
35 represents dibenzylideneacetone, "DCC" represents 1,3-dicyclohexylcarbodiimide, "DIBAH" represents diisobutylaluminum hydride, "DIPEA" represents diisopropylethylamine, "DMF" represents

dimethylformamide, "DMSO" represents dimethylsulfoxide. "Fmoc" represents 9-fluorenylmethoxycarbonyl. "LDA" represents lithium diisopropylamide, "PHTH" represents a phthaloyl group, "pnZ" represents 4-nitrobenzyloxycarbonyl, "PTC" represents a phase transfer catalyst, "p-TsOH" represents paratoluenesulfonic acid, "TBAF" represents tetrabutylammonium fluoride, "TBTU" represents 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate, "TEA" represents triethylamine, "TFA" represents trifluoroacetic acid, "THF" represents tetrahydrofuran, "TMS" represents trimethylsilyl, and "Z" represents benzyloxycarbonyl.

The present invention comprises a pharmaceutical composition comprising a therapeutically-effective amount of a compound of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP in association with at least one pharmaceutically-acceptable carrier, adjuvant or diluent.

The present invention also comprises a treatment and prophylaxis of coronary artery disease and other CETP-mediated disorders in a subject, comprising administering to the subject having such disorder a therapeutically-effective amount of a compound of Formula I-H:



(I-H)

or a pharmaceutically-acceptable salt thereof, wherein R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} , R_{12} , R_{13} , R_{14} , R_{15} , R_{16} , X, Y, and Z are as defined above for the compounds of Formula I-H.

As a further embodiment, compounds of the present invention of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP or a pharmaceutically-acceptable salt thereof as defined above comprise a treatment and prophylaxis of coronary artery disease and other CETP-mediated disorders in a subject, comprising administering to the subject having such disorder a therapeutically-effective amount of compounds I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP of the present invention or a pharmaceutically-acceptable salt thereof.

Compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP are capable of inhibiting activity of cholesteryl ester transfer protein (CETP), and thus could be used in the manufacture of a medicament, a method for the prophylactic or therapeutic treatment of diseases mediated by CETP, such as peripheral vascular disease, hyperlipidaemia, hypercholesterolemia, and other diseases attributable to either high LDL and low HDL or a combination of both, or a procedure to study the mechanism of action of the cholesteryl ester transfer protein (CETP) to enable the design of better inhibitors. The compounds of Formula I-H would be also useful in prevention of cerebral vascular accident (CVA) or stroke.

Also included in the family of compounds of Formula I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP are the pharmaceutically-acceptable salts thereof. The term "pharmaceutically-acceptable salts" embraces salts commonly used to form alkali metal salts and to form addition salts of free acids or free bases. The nature of the salt is not critical, provided that it is pharmaceutically acceptable. Suitable pharmaceutically-acceptable acid addition salts of compounds of Formula I-H may be prepared from inorganic acid or from an organic acid. Examples of such inorganic acids are hydrochloric, hydrobromic, hydroiodic, nitric, carbonic, sulfuric and phosphoric acid. Appropriate organic acids may be selected from aliphatic, cycloaliphatic, aromatic, araliphatic, heterocyclic, carboxylic and sulfonic classes of organic acids, examples of which are formic, acetic, propionic, succinic, glycolic, gluconic, lactic, malic, tartaric, citric, ascorbic, glucuronic, maleic, fumaric, pyruvic, aspartic, glutamic, benzoic, anthranilic, mesylic, salicylic, p-hydroxybenzoic, phenylacetic, mandelic, embonic (pamoic), methanesulfonic, ethylsulfonic, benzenesulfonic, sulfanilic, stearic, cyclohexylaminosulfonic, algenic, galacturonic acid. Suitable pharmaceutically-acceptable base addition salts of compounds of Formula V-H include metallic salts made from aluminum, calcium, lithium, magnesium, potassium, sodium and zinc or

organic salts made from N,N'-dibenzylethylenediamine, choline, chloroprocaine, diethanolamine, ethylenediamine, meglumine (N-methylglucamine) and procain. All of these salts may be prepared by conventional means from the corresponding compound of Formula I-H by
5 reacting, for example, the appropriate acid or base with the compound of Formula I-H.

Also embraced within this invention is a class of pharmaceutical compositions comprising the active compounds of Formula I-H in association with one or more non-toxic, pharmaceutically-acceptable carriers and/or
10 diluents and/or adjuvants (collectively referred to herein as "carrier" materials) and, if desired, other active ingredients. The active compounds of the present invention may be administered by any suitable route, preferably in the form of a pharmaceutical composition adapted to such a route, and in a dose effective for the treatment intended. The active compounds and composition may, for
15 example, be administered orally, intravascularly, intraperitoneally, subcutaneously, intramuscularly or topically.

For oral administration, the pharmaceutical composition may be in the form of, for example, a tablet, capsule, suspension or liquid. The pharmaceutical composition is preferably made in the form of a dosage unit
20 containing a particular amount of the active ingredient. Examples of such dosage units are tablets or capsules. The active ingredient may also be administered by injection as a composition wherein, for example, saline, dextrose or water may be used as a suitable carrier.

The amount of therapeutically active compounds which are
25 administered and the dosage regimen for treating a disease condition with the compounds and/or compositions of this invention depends on a variety of factors, including the age, weight, sex and medical condition of the subject, the severity of the disease, the route and frequency of administration, and the particular compound employed, and thus may vary widely.

30 The pharmaceutical compositions may contain active ingredients in the range of about 0.1 to 2000 mg, and preferably in the range of about 0.5 to 500 mg. A daily dose of about 0.01 to 100 mg/kg body weight, and preferably between about 0.5 and about 20 mg/kg body weight, may be appropriate. The daily dose can be administered in one to four doses per day.

35 The compounds may be formulated in topical ointment or cream, or as a suppository, containing the active ingredients in a total amount of, for example, 0.075 to 30% w/w, preferably 0.2 to 20% w/w and most preferably 0.4 to

15% w/w. When formulated in an ointment, the active ingredients may be employed with either paraffinic or a water-miscible ointment base.

Alternatively, the active ingredients may be formulated in a cream with an oil-in-water cream base. If desired, the aqueous phase of the cream base may
5 include, for example at least 30% w/w of a polyhydric alcohol such as propylene glycol, butane-1,3-diol, mannitol, sorbitol, glycerol, polyethylene glycol and mixtures thereof. The topical formulation may desirably include a compound which enhances absorption or penetration of the active ingredient through the skin or other affected areas. Examples of such dermal penetration
10 enhancers include dimethylsulfoxide and related analogs. The compounds of this invention can also be administered by a transdermal device. Preferably topical administration will be accomplished using a patch either of the reservoir and porous membrane type or of a solid matrix variety. In either case, the active agent is delivered continuously from the reservoir or microcapsules
15 through a membrane into the active agent permeable adhesive, which is in contact with the skin or mucosa of the recipient. If the active agent is absorbed through the skin, a controlled and predetermined flow of the active agent is administered to the recipient. In the case of microcapsules, the encapsulating agent may also function as the membrane.

20 The oily phase of the emulsions of this invention may be constituted from known ingredients in a known manner. While the phase may comprise merely an emulsifier, it may comprise a mixture of at least one emulsifier with a fat or an oil or with both a fat and an oil. Preferably, a hydrophilic emulsifier is included together with a lipophilic emulsifier which acts as a stabilizer. It is
25 also preferred to include both an oil and a fat. Together, the emulsifier(s) with or without stabilizer(s) make-up the so-called emulsifying wax, and the wax together with the oil and fat make up the so-called emulsifying ointment base which forms the oily dispersed phase of the cream formulations. Emulsifiers and emulsion stabilizers suitable for use in the formulation of the present
30 invention include Tween 60, Span 80, cetostearyl alcohol, myristyl alcohol, glyceryl monostearate, and sodium lauryl sulfate, among others.

The choice of suitable oils or fats for the formulation is based on achieving the desired cosmetic properties, since the solubility of the active compound in most oils likely to be used in pharmaceutical emulsion
35 formulations is very low. Thus, the cream should preferably be a non-greasy, non-staining and washable product with suitable consistency to avoid leakage from tubes or other containers. Straight or branched chain, mono- or dibasic

alkyl esters such as di-isoadipate, isocetyl stearate, propylene glycol diester of coconut fatty acids, isopropyl myristate, decyl oleate, isopropyl palmitate, butyl stearate, 2-ethylhexyl palmitate or a blend of branched chain esters may be used. These may be used alone or in combination depending on the
5 properties required. Alternatively, high melting point lipids such as white soft paraffin and/or liquid paraffin or other mineral oils can be used.

For therapeutic purposes, the active compounds of this combination invention are ordinarily combined with one or more adjuvants appropriate to the indicated route of administration. If administered *per os*, the compounds
10 may be admixed with lactose, sucrose, starch powder, cellulose esters of alkanolic acids, cellulose alkyl esters, talc, stearic acid, magnesium stearate, magnesium oxide, sodium and calcium salts of phosphoric and sulfuric acids, gelatin, acacia gum, sodium alginate, polyvinylpyrrolidone, and/or polyvinyl alcohol, and then tableted or encapsulated for convenient administration. Such
15 capsules or tablets may contain a controlled-release formulation as may be provided in a dispersion of active compound in hydroxypropylmethyl cellulose. Formulations for parenteral administration may be in the form of aqueous or non-aqueous isotonic sterile injection solutions or suspensions. These solutions and suspensions may be prepared from sterile powders or
20 granules having one or more of the carriers or diluents mentioned for use in the formulations for oral administration. The compounds may be dissolved in water, polyethylene glycol, propylene glycol, ethanol, corn oil, cottonseed oil, peanut oil, sesame oil, benzyl alcohol, sodium chloride, and/or various buffers. Other adjuvants and modes of administration are well and widely
25 known in the pharmaceutical art.

The present invention further comprises a process for the preparation of (R)-chiral compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP by reacting suitable secondary amines with (R)-chiral forms of alcohols, epoxides, and cyclic sulfate esters.

30 The present invention also comprises a process for the preparation of (R)-chiral compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP by reacting a suitable secondary amine with a substantially stoichiometric amount of a (R)-chiral epoxide in the presence of a transition metal-based salt.

35 The present invention also comprises a process for the preparation of (R)-chiral precursor compounds useful in the preparation of compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-

CP by reacting a suitable primary amine with a substantially stoichiometric amount of a (R)-chiral epoxide with or without the presence of an added transition metal-based compound.

5 All mentioned references are incorporated by reference as if here written.

Although this invention has been described with respect to specific embodiments, the details of these embodiments are not to be construed as limitations.

10 GENERAL SYNTHETIC PROCEDURES

The compounds of the present invention can be synthesized, for example, according to the following procedures of Schemes 1 through 15 below, wherein the substituents are as defined for Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP above except where further noted.

Synthetic Scheme 1 shows the preparation of compounds of formula XIII A-H ("Secondary Heteroaryl Amines") which are intermediates in the preparation of the compounds of the present invention corresponding to
20 Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP
25 ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") wherein A and Q are independently aryl and heteroaryl. Schemes 1 through 3, taken together, prepare 1-substitutedamino-2-alkanols of the present invention by addition of a halogenated, oxygen containing precursor to a secondary
30 amine to introduce an oxy containing alkyl group wherein the two groups making up the secondary amine both are made up of aromatic groups or both groups contain aromatic rings wherein said aromatic rings maybe 0 to 2 aryl rings and 0 to 2 heteroaryl rings.

The "Generic Imine" corresponding to Formula XII can be prepared
35 through dehydration techniques generally known in or adaptable from the art by reacting "Generic Amine-I" of Formula X with the "Generic Carbonyl

Compound" of Formula XI in Scheme 1 and subsequent specific examples. For example, when Z is a covalent bond, methylene, methine substituted with another substituent, ethylene, or another substituent as defined in Formula I-H, the two reactants (X and XI) react by refluxing them in an aprotic solvent, such as hexane, toluene, cyclohexane, benzene, and the like, using a Dean-Stark type trap to remove water. After about 2-8 hours or until the removal of water is complete, the aprotic solvent is removed *in vacuo* to yield the "Generic Imine" of Formula XII. Alternately, when Z is an oxygen, the "Generic Imine" is an oxime derivative. Alternately, when Z is a nitrogen, the "Generic Imine" is a hydrazone derivative. Hydrazone type "Generic Imine" compounds are readily prepared from the corresponding hydrazine and the appropriate aldehyde or ketone type "Generic Carbonyl Compound". Suitable procedures for forming oxime and hydrazone imines are also described by Shriner, Fuson, and Curtin in *The Systematic Identification of Organic Compounds*, 5th Edition, John Wiley & Sons, and by Fieser and Fieser in *Reagents for Organic Synthesis*, Volume 1, John Wiley & Sons, which are incorporated herein by reference.

The "Generic Secondary Amines" of Formula XIII can be prepared from the corresponding "Generic Imine" of Formula XII in several ways. For example, in one synthetic scheme (Reduction Method-1), which is preferred when Z is a nitrogen, the "Generic Imine" hydrazone of Formula XII is partially or completely dissolved in lower alkanols containing sufficient organic acid or mineral acid as described in WO Patent Application No.9738973, Swiss Patent CH 441366 and U. S. Patent Nos. 3359316 and 3334017, which are incorporated herein by reference, and then hydrogenated at 0-100°C, more preferably 20-50°C, and most preferably between 20-30°C and pressures of 10-200 psi hydrogen or more preferably between 50-70 psi hydrogen in the presence of a noble metal catalyst such as PtO₂.

In another synthetic scheme (Reduction Method-2), which is preferred when Z is a single bond or carbon, the "Generic Imine" of Formula XII is slurried in a lower alcohol such as ethanol, methanol or like solvent at 0-10°C and solid sodium borohydride is added in batches over 5-10 minutes at 0-10°C with stirring. The reaction mixture is stirred below 10°C for 30-90 minutes

and then is warmed gradually to 15-30°C. After about 1-10 hours, the mixture is cooled and acid is added until the aqueous layer was just acidic (pH 5-7).

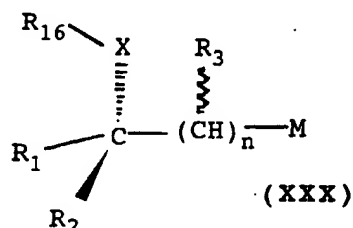
In yet another synthetic scheme (Reduction Method-3), which is preferred when Z is an oxygen, the "Generic Imine" oxime of Formula XII is
5 slurried in a lower alcohol solvent such methanol or like solvent at 0-10°C and acidified to a pH less than 4. Solid sodium cyanoborohydride is added in batches over 30-90 minutes at 0-20°C with stirring and addition of a suitable organic or mineral acid to keep the pH at or below 4. The reaction mixture is stirred and warmed gradually to about 20-25°C. After about 1-10 hours, the
10 mixture is cooled and base added until the mixture was just slightly alkaline.

The "Generic Secondary Amines" of Formula XIII can also be prepared, according to Scheme 1 by an alkylation procedures based on the nucleophilic substitution of bromides by amines. In one synthetic alkylation scheme (Alkylation Method-1), a "Generic Amine-1" of Formula X is reacted
15 with a "Generic Bromide-2" of Formula XXIII as described in Vogel's Textbook of Practical Organic Chemistry, Fifth Edition, 1989, pages 902 to 905 and references cited therein all of which are incorporated herein by reference. In an alternate synthetic alkylation scheme (Alkylation Method-2), a "Generic Amine-2" of Formula XXII is reacted with a "Generic Bromide-2" of
20 Formula XXIII in a method employing palladium catalyzed carbon-nitrogen bond formation. Suitable procedures for this conversion are described in Wagaw and Buchwald, J. Org. Chem.(1996), 61, 7240-7241, Wolfe, Wagaw and Buchwald, J. Am. Chem. Soc. (1996), 118, 7215-7216, and Wolfe and Buchwald, Tetrahedron Letters (1997), 38(36), 6359-6362 and references
25 cited therein all of which are incorporated herein by reference. The preferred "Generic Bromide-2" of Formula XXIII are generally aryl bromides, aryl triflates, and heteroaryl bromides.

The "Generic Amine-1" and "Generic Amine-2" amines, hydroxylamines, and hydrazines, the "Generic Carbonyl Compound"
30 aldehydes, ketones, hydrazones, and oximes, and "Generic Bromide-1" and "Generic Bromide-2" halides, tosylates, mesylates, triflates, and precursor alcohols required to prepare the "Generic Secondary Amine" compounds are available from commercial sources or can be prepared by one skilled in the art from published procedures. Commercial sources include but are not limited to
35 Aldrich Chemical, TCI-America, Lancaster-Synthesis, Oakwood Products,

- Acros Organics, and Maybridge Chemical. Disclosed procedures for "Generic Amine" amines, hydroxylamines, and hydrazines include Sheradsky and Nov. J. Chem. Soc., Perkin Trans.1 (1980), (12), 2781-6; Marcoux, Doye, and Buchwald, J. Am. Chem. Soc. (1997), 119, 1053-9; Sternbach and Jamison, 5 Tetrahedron Lett. (1981), 22(35), 3331-4; U. S. Patent No. 5306718; EP No. 314435; WO No. 9001874; WO No. 9002113; JP No. 05320117; WO No. 9738973; Swiss Patent No. CH 441366; U. S. Patents Nos. 3359316 and 3334017; and references cited therein which are incorporated herein by reference.
- 10 Synthetic Scheme 2 shows the preparation of the class of compounds of the present invention corresponding to Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic 15 Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") wherein A and Q are independently aryl and heteroaryl.
- 20 Derivatives of "Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols", "Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols", "Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols", "Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-25 (n+1)-Alkanols", and "Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols", in which the hetero atom (-O-) is attached to an alkyl group removed from the amine by two or more carbons are readily prepared by anion chemistry using the method of Scheme 2. The anion of "Generic Secondary Amine" amines, hydroxylamines, and hydrazines of 30 Formula XIII is readily formed by dissolving the specific amine, hydroxylamine, or hydrazine in an aprotic solvent, such as tetrahydrofuran, toluene, ether, dimethylformamide, and dimethylformamide, under anhydrous conditions. The solution is cooled to a temperature between -78 and 0°C, preferably between -78 and -60°C, and the anion formed by the addition of at 35 least one equivalent of a strong, aprotic, non-nucleophilic base, such as NaH or n-butyllithium under an inert atmosphere, for each acidic group present.

Maintaining the temperature between -78 and 0°C, preferably between -78 and -60°C, with suitable cooling, an appropriate alkyl halide, alkyl benzenesulfonate such as a alkyl tosylate, alkyl mesylate, alkyl triflate or similar alkylating reagent of the general structure:



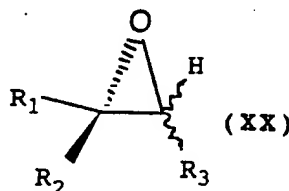
where M is a readily displaceable group such as chloride, bromide, iodide, tosylate, triflate, and mesylate and X is oxy. After allowing the reaction mixture to warm to room temperature, the reaction product is added to water, neutralized if necessary, and extracted with a water-immiscible solvent such as diethyl ether or methylene chloride. The combined aprotic solvent extract is washed with saturated brine, dried over drying agent such as anhydrous MgSO₄ and concentrated *in vacuo* to yield crude Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"). This material is purified, for example, by eluting through silica gel with a medium polar solvent such as ethyl acetate in a non-polar solvent such as hexanes to yield Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"). Products are structurally confirmed by low and high resolution mass spectrometry and NMR.

Compounds of Formula (XXX), which can be used to prepare the "Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanol" compounds of Tables 3 and 4, are given in Table 2. Reagents 1a and 2a in Table 2 are prepared from the corresponding alcohols. (R)-Chiral alcohol precursors to 1a, 2a, and similar alcohols that can be envisioned by one of inventive skill can be obtained from the corresponding racemic mixture of the R-enantiomer and S-enantiomer by separation procedures using preparative gas chromatography and high pressure liquid chromatography using chiral chromatographic columns. The tosylates of chiral alcohols and racemic mixtures are readily obtained by reacting the corresponding alcohol with tosyl chloride using procedures found in House's Modern Synthetic Reactions, Chapter 7, W. A. Benjamin, Inc., Shriner, Fuson, and Curtin in The Systematic Identification of Organic Compounds, 5th Edition, John Wiley & Sons, and Fieser and Fieser in Reagents for Organic Synthesis, Volume 1, John Wiley & Sons, which are incorporated herein by reference.

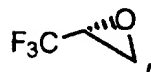
Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds can also be prepared using the method of Scheme 2 through the use of racemic (XXX) as described followed by preparative separation of the R-enantiomer from the S-enantiomer using chiral chromatographic procedures such as preparative gas chromatography and high pressure liquid chromatography using readily available chiral chromatographic columns and procedures.

A preferred procedure for Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds is the novel inventive Method A of Scheme 3. (R)-Chiral oxirane reagents useful in Method A are exemplified, but not limited to those in Table 1. Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl

(R)-Chiral Halogenated 1-Substitutedamino-2-Propanols”), and Formula I-CP (“Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols”) compounds are prepared by reacting “Generic Secondary Amine” amines, hydroxylamines, and hydrazines of Formula XIII with (R)-chiral oxiranes of
 5 the type listed in Table 1 and represented by the general structure:



Oxiranes having a specific stereochemical arrangement of R_1 , R_2 and R_3 can be prepared using chiral procedures such as those published in 1995 by Ramachandran, Gong, and Brown in the Journal of Organic Chemistry, Vol. 60,
 10 pages 41 to 46; cited references also detail alternate procedures to prepare chiral and achiral epoxides, which are incorporated herein by reference. For example, the specific preparation of *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane.

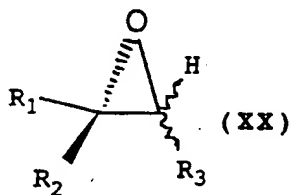


, using a procedure adopted from H.C.Brown et al. (*J. Org. Chem.* 60, 41-46, (1995)), is accomplished as described in **Example 4**. Many
 15 of the epoxides summarized in Table 1 can be prepared in the (R)-configuration using procedures analogous to that given above for *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane.

In some cases, achiral oxiranes of (XX) can be prepared from the corresponding alkenes by reaction of epoxidation reagents such as meta-chloroperbenzoic acid (MCPBA) and similar type reagents readily selectable by
 20 a person of skill-in-the-art with alkenes. Fieser and Fieser in Reagents for Organic Synthesis, John Wiley & Sons provides, along with cited references, numerous suitable epoxidation reagents and reaction conditions, which are incorporated herein by reference. These achiral oxiranes can be reacted in an
 25 identical manner to that described for (R)-chiral oxiranes with “Generic Secondary Amine” amines, hydroxylamines, and hydrazines of Formula XIII to afford racemic compounds structurally identical to those of Formula I-HP, Formula I-HPC, and Formula I-C but with the corresponding (S) chiral configuration present in an equivalent amount. Formula I-HP (“Generic

- Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols”), Formula I-HPC (“Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols”), and Formula I-CP (“Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols”)
- 5 compounds can be obtained by preparative chiral chromatography of said racemic mixtures to obtain the (R)-chiral configuration of Formula I-HP, Formula I-HPC, and Formula I-CP substantially free of the (S)-chiral configuration enantiomer. Alternatively, achiral oxiranes may be separated by chiral preparative chromatography into their respective (R)-Chiral and (S)-
- 10 Chiral enantiomers and the (R)-Chiral enantiomer reacted to afford Formula I-HP (“Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols”), Formula I-HPC (“Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols”), and Formula I-CP (“Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols”)
- 15 compounds.

Table 1. Structure of (R)-Chiral Oxirane Reagents.

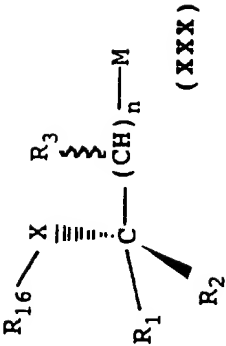


| Reagent Number | <u>R₁</u> | <u>R₂</u> | <u>R₃</u> |
|-------------------|--|------------------------------------|----------------------|
| 1 | CF ₃ | H | H |
| 2 | CCl ₃ | H | H |
| 3 | CF ₃ | CH ₃ | H |
| 4 | CF ₃ CF ₂ | H | H |
| 5 | CF ₃ CF ₂ CF ₂ | H | H |
| 6 | CF ₃ OCF ₂ CF ₂ | H | H |
| 7 | CF ₃ CH ₂ | H | H |
| 9 | CF ₃ | H | CF ₃ |
| 11 | CF ₃ | C ₆ H ₅ | H |
| 12 | CCl ₃ | C ₆ H ₅ | H |
| 13 | CCl ₃ | Cyclopropyl | H |
| 14 | CCl ₃ | CH ₃ | H |
| 15 | CCl ₃ | (CH ₃) ₂ CH | H |
| 16 | CHCl ₂ | H | H |
| 18 | CF ₃ | H | CH ₃ |
| 27 | CCl ₃ CH ₂ | H | H |
| 28 | CBr ₃ CH ₂ | H | H |
| 29 | CHBr ₂ CH ₂ | H | H |
| 30 | CBrCl ₂ | H | H |
| 31 | CClF ₂ | H | H |
| 32 | CCl ₂ F | H | H |

Table 1. (continued) Structure of (R)-Chiral Oxirane Reagents.

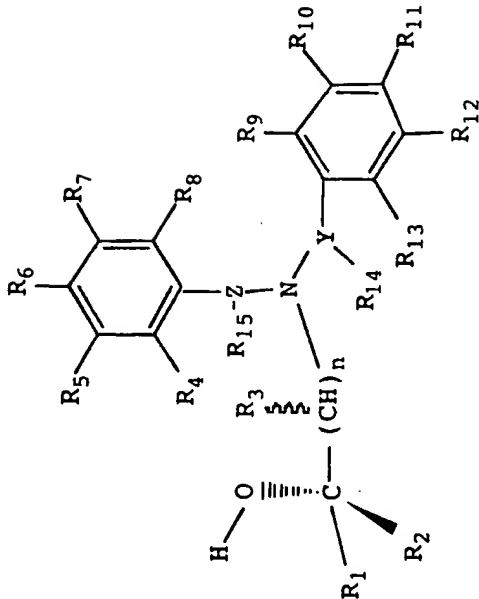
| <u>Reagent Number</u> | <u>R₁</u> | <u>R₂</u> | <u>R₃</u> |
|---------------------------|---|----------------------|----------------------|
| 33 | CCl ₃ CCl ₂ | H | H |
| 43 | FCH ₂ | H | H |
| 56 | CBrF ₂ CClFCH ₂ | H | H |
| 57 | HCF ₂ CF ₂ OCH ₂ | H | H |

Table 2. Structure and Source of Alcohol and Glycol Reagents.



| Reagent Number | <u>R₁</u> | <u>n</u> | <u>M</u> | <u>R₂</u> | <u>R₃</u> | <u>X-R₁₆</u> | Source of Reagent |
|----------------|---|----------|----------|----------------------|----------------------|-------------------------|--|
| 1A | CF ₃ | 3 | OTs | H | H | OH | Chiral separation and then tosylation of alcohol from Justus Liebigs Ann. Chem. (1969), 720, 81-97. |
| 2A | CF ₃ CH ₂ CH ₂ | 3 | OTs | H | H | OH | Chiral separation and then tosylation of alcohol from Z. Naturforsch., B: Chem. Sci. (1997), 52 (3), 413-418 |

Table 3. Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).



| Inhibitor Number Column 1 + Column 2 | | R ₁ | R ₂ | R ₃ | R ₄ | R ₅ | R ₆ | R ₇ | R ₁₀ | R ₁₁ |
|---|---------|-----------------|----------------|----------------|----------------|---------------------------------|----------------|----------------|------------------------------------|-----------------|
| Reagent | Reagent | | | | | | | | | |
| 1A | 1N | CF ₃ | H | H | H | C ₆ H ₅ O | H | H | OCF ₂ CF ₂ H | H |
| 1A | 2N | CF ₃ | H | H | H | OCF ₃ | H | H | OCF ₂ CF ₂ H | H |

Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| <u>Inhibitor Number</u> <u>Column 1 + Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₇</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|---------------------------------|----------------------|----------------------|------------------------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| 1A | 3N | CF ₃ | 3 | H | H | F | H | H | F | OCF ₂ CF ₂ H | H |
| 1A | 4N | CF ₃ | 3 | H | H | H | F | H | H | OCF ₂ CF ₂ H | H |
| 1A | 5N | CF ₃ | 3 | H | H | H | C ₆ H ₅ O | H | H | OCF ₃ | H |
| 1A | 6N | CF ₃ | 3 | H | H | H | OCF ₃ | H | H | OCF ₃ | H |
| 1A | 7N | CF ₃ | 3 | H | H | H | H | phenyl | H | OCF ₃ | H |
| 1A | 8N | CF ₃ | 3 | H | H | H | phenyl | H | H | OCF ₃ | H |
| 1A | 9N | CF ₃ | 3 | H | H | H | H | H | H | OCF ₃ | H |
| 1A | 10N | CF ₃ | 3 | H | H | H | Br | H | H | OCF ₃ | H |
| 1A | 11N | CF ₃ | 3 | H | H | H | CF ₃ | F | H | CF ₃ | H |
| 1A | 12N | CF ₃ | 3 | H | H | H | CH ₃ | H | H | CF ₃ | H |
| 1A | 13N | CF ₃ | 3 | H | H | H | CF ₃ | H | H | CF ₃ | H |
| 1A | 14N | CF ₃ | 3 | H | H | H | CH ₃ | H | H | OCF ₃ | H |

Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| Inhibitor Number Column1+Column 2 | | R ₁ | n | R ₂ | R ₃ | R ₄ | R ₅ | R ₆ | R ₇ | R ₁₀ | R ₁₁ |
|--------------------------------------|---------|-----------------|---|----------------|----------------|----------------|-----------------|----------------|----------------|------------------|-----------------|
| Reagent | Reagent | | | | | | | | | | |
| 1A | 15N | CF ₃ | 3 | H | H | H | F | F | | | |
| 1A | 16N | CF ₃ | 3 | H | H | H | Br | H | H | OCF ₃ | H |
| 1A | 17N | CF ₃ | 3 | H | H | H | CF ₃ | F | H | OCF ₃ | H |
| 1A | 18N | CF ₃ | 3 | H | H | H | F | H | H | OCF ₃ | H |
| 1A | 19N | CF ₃ | 3 | H | H | H | Cl | H | H | OCF ₃ | H |
| 1A | 20N | CF ₃ | 3 | H | H | H | F | H | H | CF ₃ | H |
| 1A | 21N | CF ₃ | 3 | H | H | H | F | F | H | CF ₃ | H |
| 1A | 22N | CF ₃ | 3 | H | H | H | Cl | H | H | CF ₃ | H |
| 1A | 23N | CF ₃ | 3 | H | H | H | F | H | H | phenoxy | H |
| 1A | 24N | CF ₃ | 3 | H | H | H | CF ₃ | Cl | H | CH ₃ | H |
| 1A | 25N | CF ₃ | 3 | H | H | H | CF ₃ | F | H | CH ₃ | H |
| 1A | 26N | CF ₃ | 3 | H | H | H | H | H | H | CF ₃ | H |

Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| <u>Inhibitor Number</u> <u>Column 1 + Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₇</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| IA | 27N | CF ₃ | 3 | H | H | F | F | H | H | CF ₃ | H |
| IA | 28N | CF ₃ | 3 | H | H | H | H | OCH ₃ | H | CF ₃ | H |
| IA | 29N | CF ₃ | 3 | H | H | H | F | F | H | CH ₃ | H |
| IA | 30N | CF ₃ | 3 | H | H | H | OCH ₃ | H | H | CH ₃ | H |
| IA | 31N | CF ₃ | 3 | H | H | H | H | CH ₃ | H | H | H |
| IA | 32N | CF ₃ | 3 | H | H | H | Cl | H | H | H | H |
| IA | 33N | CF ₃ | 3 | H | H | H | F | H | H | F | H |
| IA | 34N | CF ₃ | 3 | H | H | H | H | OCH ₃ | H | CH ₃ | H |
| IA | 35N | CF ₃ | 3 | H | H | H | H | H | H | H | H |
| IA | 36N | CF ₃ | 3 | H | H | H | H | CH ₃ | H | CH ₃ | H |
| IA | 37N | CF ₃ | 3 | H | H | H | H | Cl | H | H | H |
| IA | 38N | CF ₃ | 3 | H | H | H | F | H | H | 3-CF ₃ - | H |

Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| <u>Inhibitor Number</u> <u>Column 1 + Column 2</u> | | <u>R₁</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₇</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|---------------------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | |
| IA | 39N | CF ₃ | H | H | H | F | H | H | phenoxy | H |
| IA | 40N | CF ₃ | H | H | H | F | H | H | 4-CH ₃ O- phenoxy | H |
| IA | 41N | CF ₃ | H | H | H | F | H | H | 4-Cl- phenoxy | H |
| IA | 42N | CF ₃ | H | H | H | F | H | H | H | H |
| IA | 43N | CF ₃ | H | H | H | F | H | F | CH ₃ | H |
| IA | 44N | CF ₃ | H | H | F | F | H | H | CH ₃ | H |
| IA | 45N | CF ₃ | H | H | H | Cl | H | H | CH ₃ | H |
| IA | 46N | CF ₃ | H | H | H | CH ₃ | H | H | CH ₃ | H |
| IA | 48N | CF ₃ | H | H | H | H | CH ₃ | H | CF ₃ | H |
| IA | 51N | CF ₃ | H | H | H | H | CH ₃ | H | F | H |

Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| <u>Inhibitor Number</u> <u>Column 1+Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₇</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|---------------------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| IA | 52N | CF ₃ | 3 | H | H | H | CF ₃ | H | H | F | H |
| IA | 53N | CF ₃ | 3 | H | H | H | CF ₃ | H | H | CH ₃ | H |
| IA | 54N | CF ₃ | 3 | H | H | H | OCH ₃ | H | H | CF ₃ | H |
| IA | 56N | CF ₃ | 3 | H | H | H | H | CH ₃ | H | CF ₃ | H |
| IA | 57N | CF ₃ | 3 | H | H | H | C ₆ H ₅ O | H | H | H | OCF ₃ |
| IA | 58N | CF ₃ | 3 | H | H | H | H | H | H | H | OCF ₃ |
| IA | 59N | CF ₃ | 3 | H | H | H | OCF ₃ | H | H | H | OCF ₃ |
| IA | 60N | CF ₃ | 3 | H | H | H | CF ₃ | F | H | H | CF ₃ |
| IA | 61N | CF ₃ | 3 | H | H | H | H | OCH ₃ | H | H | CF ₃ |
| IA | 62N | CF ₃ | 3 | H | H | H | CH ₃ | H | H | H | CF ₃ |
| IA | 63N | CF ₃ | 3 | H | H | H | Cl | H | H | H | CF ₃ |
| IA | 64N | CF ₃ | 3 | H | H | H | CF ₃ | H | H | H | OCF ₃ |

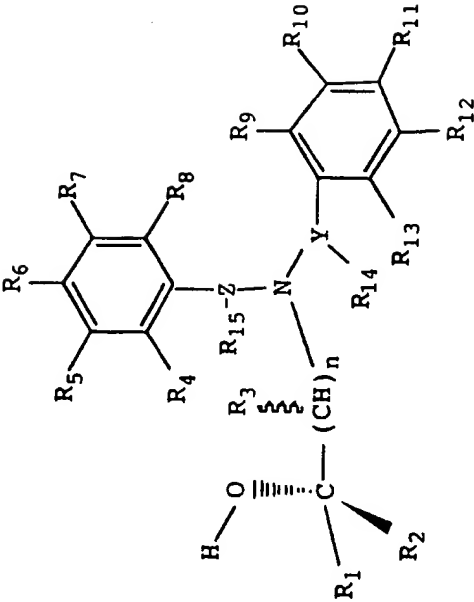
Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| <u>Inhibitor Number</u> <u>Column 1 + Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₇</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| IA | 65N | CF ₃ | 3 | H | H | H | F | H | H | H | OCF ₃ |
| IA | 66N | CF ₃ | 3 | H | H | H | F | H | F | H | OCF ₃ |
| IA | 67N | CF ₃ | 3 | H | H | H | Br | H | H | H | OCF ₃ |
| IA | 68N | CF ₃ | 3 | H | H | H | Cl | H | H | H | OCF ₃ |
| IA | 69N | CF ₃ | 3 | H | H | H | F | F | H | H | OCF ₃ |
| IA | 70N | CF ₃ | 3 | H | H | H | F | H | H | H | OCF ₃ |
| IA | 71N | CF ₃ | 3 | H | H | H | CH ₃ | H | H | H | phenyl |
| IA | 72N | CF ₃ | 3 | H | H | H | F | F | H | H | OCF ₃ |
| IA | 73N | CF ₃ | 3 | H | H | H | Cl | H | H | H | CF ₃ |
| IA | 74N | CF ₃ | 3 | H | H | H | OCH ₃ | H | H | H | CH ₃ |
| IA | 75N | CF ₃ | 3 | H | H | H | F | H | H | H | CH ₃ |
| IA | 76N | CF ₃ | 3 | H | H | H | F | H | H | H | OCF ₃ |

Table 3. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y is CH; R₈, R₉, R₁₂, R₁₃, and R₁₄ are each H; Z is covalent bond and R₁₅ is absent).

| Inhibitor Number Column 1 + Column 2 | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₇</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| 1A | 78N | CF ₃ | 3 | H | H | H | H | OCH ₃ | H | H | CH ₃ |
| 1A | 79N | CF ₃ | 3 | H | H | H | H | CH ₃ | H | H | CH ₃ |
| 1A | 80N | CF ₃ | 3 | H | H | H | CH ₃ | H | H | H | CH ₃ |
| 1A | 82N | CF ₃ | 3 | H | H | H | F | F | H | H | CH ₃ |
| 1A | 83N | CF ₃ | 3 | H | H | H | F | H | F | H | CH ₃ |
| 1A | 84N | CF ₃ | 3 | H | H | F | F | H | H | H | CH ₃ |
| 1A | 85N | CF ₃ | 3 | H | H | F | CF ₃ | H | H | H | CH ₃ |
| 1A | 86N | CF ₃ | 3 | H | H | H | H | CH ₃ | H | H | CF ₃ |
| 1A | 88N | CF ₃ | 3 | H | H | H | CF ₃ | H | H | H | CH ₃ |
| 1A | 90N | CF ₃ | 3 | H | H | H | H | CF ₃ | H | H | CH ₃ |
| 1A | 92N | CF ₃ | 3 | H | H | H | CF ₃ | F | H | H | CH ₃ |

Table 4. Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y and Z are each CH; R₈, R₉, R₁₂, R₁₃, R₁₄ and R₁₅ are each H).



| <u>Inhibitor Number</u> | | <u>Column 1+Column 2</u> | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₉</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|-------------------------|---------|--------------------------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | | |
| IA | IDB | | CF ₃ | 3 | H | H | H | OCF ₃ | H | H | OCF ₃ | H |

Table 4. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y and Z are each CH₃; R₈, R₉, R₁₂, R₁₃, R₁₄ and R₁₅ are each H).

| <u>Inhibitor Number</u> <u>Column 1+Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₉</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| 1A | 2DB | CF ₃ | 3 | H | H | H | Cl | H | H | H | CF ₃ |
| 1A | 3DB | CF ₃ | 3 | H | H | H | Br | H | H | OCF ₃ | H |
| 1A | 4DB | CF ₃ | 3 | H | H | H | Cl | H | H | OCF ₃ | H |
| 1A | 5DB | CF ₃ | 3 | H | H | H | Cl | H | H | CF ₃ | H |
| 1A | 6DB | CF ₃ | 3 | H | H | H | H | Cl | H | CF ₃ | H |
| 1A | 7DB | CF ₃ | 3 | H | H | H | F | H | H | OCF ₃ | H |
| 1A | 8DB | CF ₃ | 3 | H | H | H | H | Cl | H | H | CF ₃ |
| 1A | 9DB | CF ₃ | 3 | H | H | H | F | H | H | H | CF ₃ |
| 1A | 10DB | CF ₃ | 3 | H | H | H | H | F | H | H | CF ₃ |
| 1A | 11DB | CF ₃ | 3 | H | H | H | H | H | H | H | CF ₃ |
| 1A | 12DB | CF ₃ | 3 | H | H | H | Cl | H | CF ₃ | H | H |

Table 4. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y and Z are each CH; R₈, R₉, R₁₂, R₁₃, R₁₄ and R₁₅ are each H).

| <u>Inhibitor Number</u> <u>Column 1+Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₉</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| IA | 13DB | CF ₃ | 3 | H | H | H | H | Cl | CF ₃ | H | H |
| IA | 14DB | CF ₃ | 3 | H | H | Cl | H | H | CF ₃ | H | H |
| IA | 15DB | CF ₃ | 3 | H | H | H | F | H | CH ₃ | H | H |
| IA | 16DB | CF ₃ | 3 | H | H | H | H | F | H | H | CH ₃ |
| IA | 17DB | CF ₃ | 3 | H | H | H | F | H | H | CH ₃ | H |
| IA | 18DB | CF ₃ | 3 | H | H | F | H | H | CH ₃ | H | H |
| IA | 19DB | CF ₃ | 3 | H | H | H | H | F | H | CH ₃ | H |
| IA | 20DB | CF ₃ | 3 | H | H | F | H | H | H | H | CH ₃ |
| IA | 21DB | CF ₃ | 3 | H | H | F | H | H | H | CF ₃ | H |
| IA | 22DB | CF ₃ | 3 | H | H | Cl | H | H | H | CF ₃ | H |
| IA | 23DB | CF ₃ | 3 | H | H | H | F | H | CF ₃ | H | H |

Table 4. (continued) Structure of Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols
(Y and Z are each CH; R₈, R₉, R₁₂, R₁₃, R₁₄ and R₁₅ are each H).

| <u>Inhibitor Number</u> <u>Column 1+Column 2</u> | | <u>R₁</u> | <u>n</u> | <u>R₂</u> | <u>R₃</u> | <u>R₄</u> | <u>R₅</u> | <u>R₆</u> | <u>R₉</u> | <u>R₁₀</u> | <u>R₁₁</u> |
|---|---------|----------------------|----------|----------------------|----------------------|----------------------|----------------------|----------------------|----------------------|-----------------------|-----------------------|
| Reagent | Reagent | | | | | | | | | | |
| 1A | 24DB | CF ₃ | 3 | H | H | H | H | F | CF ₃ | H | H |
| 1A | 25DB | CF ₃ | 3 | H | H | H | F | H | H | CF ₃ | H |
| 1A | 26DB | CF ₃ | 3 | H | H | H | H | F | H | CF ₃ | H |
| 1A | 27DB | CF ₃ | 3 | H | H | H | OCF ₃ | H | H | H | OCF ₃ |

- A mixture of a "Generic Secondary Amine" amine, hydroxylamine, or hydrazine of Formula XIII and an excess of a halogenated oxirane of (R)-chiral configuration of Formula XX are stirred and heated to 40-90°C for 5 to 48 hours in a tightly capped or contained reaction vessel. More preferably, a
- 5 Lewis acid such as a transition metal-based salts (for example, ytterbium triflate, hafnium triflate, scandium triflate, neodymium triflate, gadolinium triflate, and zirconium triflate) in methylene chloride, tetrahydrofuran, or, more preferably, acetonitrile is added to speed up the reaction to a total time of 4 to 18 hours, improve yields, to permit the reaction temperature to be reduced to
- 10 15-65°C, and to use a smaller excess of halogenated oxirane. When a Lewis acid is used, the reaction should be carried out under inert, anhydrous conditions using a blanket of dry nitrogen or argon gas. After cooling to room temperature and testing the reaction mixture for complete reaction by thin layer chromatography or high pressure liquid chromatography (hplc), the reaction
- 15 product is added to water and extracted with a water immiscible solvent such as diethyl ether or methylene chloride. (Note: If the above analysis indicates that reaction is incomplete, heating should be resumed until complete with the optional addition of more of the oxirane). The combined aprotic solvent extract is washed with saturated brine, dried over drying agent such as anhydrous
- 20 MgSO_4 and concentrated *in vacuo* to yield crude Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds.
- 25 This material is purified by eluting through silica gel with 5-40% of a medium polar solvent such as ethyl acetate in a non-polar solvent such as hexanes to yield Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"),
- 30 and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds. Products are tested for purity by HPLC. If necessary, the Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-
- 35 Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds are purified

by additional chromatography or recrystallization. Products are structurally confirmed by low and high resolution mass spectrometry and NMR. Examples of specific Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds prepared are summarized in the Examples 1 through 44, and Example Tables 1 through 12.

Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds can further be prepared in an alternate manner to procedures disclosed above and in Schemes 1, 2, and 3. Schemes 9 and 10 detail such procedures to prepare aminopropanol compounds of the present invention by initial formation of an halogenated, oxygen containing primary alkylamine XVL ("Generic Substituted Alkylamine"). Said halogenated, oxygen containing primary alkylamine XVL, formed in Scheme 9, is itself converted to secondary amines, VLX-H ("Heteroaryl Alkyl Amine) and VLX ("Phenyl Alkyl Amine"), using procedures disclosed above. Primary alkylamine XVL is first reacted with an aldehydic or ketonic carbonyl compound, XI-AH ("Heteroaryl Carbonyl") with azeotropic distillation to form imine, VL-H ("Heteroaryl Imine"). Said imine VL-H is then reduced with or without prior isolation by Reduction Methods 1, 2 or 3 as disclosed above and in Scheme 1 to yield secondary amine, VLX-H ("Heteroaryl Alkyl Amine). Said secondary amine VLX-H can be converted according to Scheme 10 to give Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols") and Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds. Using similar Schemes, VLX can be converted to Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds. Compounds of this invention in which one aromatic substituent is aryl and the other aromatic substituent is heteroaryl can be readily prepared by reacting VLX-H with an aralkyl bromide or aryl bromide instead of using an heteroaralkyl bromide or heteroaryl bromide. Similarly, compounds of this

invention in which one aromatic substituent is aryl and the other aromatic substituent is heteroaryl can be readily prepared by reacting VLX with an heteroaryl bromide or heteroaralkyl bromide instead of using an aryl bromide or an aralkyl bromide.

5 Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds can further be prepared in an
10 alternate manner to procedures disclosed above and in Schemes 1, 2, 3, 9, and 10. Schemes 13, 14, and 15 detail alternate procedures to prepare (R)-Chiral Halogenated 1-Substitutedamino-2-propanols" compounds of the present invention by initial formation of an halogenated, oxygen containing secondary
15 alkylamines VLX and VLXX ("Phenyl Alkylamines") and VLXX-O ("Phenyl Oxy Alkylamines"). Said secondary alkylamines VLX and VLXX ("Phenyl Alkylamines") and VLXX-O ("Phenyl Oxy Alkylamines") can be converted according to Schemes 13, 14, and 15 to Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-
20 propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with appropriate aromatic halides such as aryl bromides and heteroaryl bromides as desired.

 Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral
25 Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds can further be prepared in an alternate manner to procedures disclosed above and in Schemes 1, 2, 3, 9, 10,
30 13, 14, and 15. Another alternate procedure to prepare "(R)-Chiral Halogenated 1-Substitutedamino-2-propanols" compounds of the present invention can be achieved by reacting secondary amines of Formula XIII A-H ("Secondary Heteroaryl Amines") and Formula XIII-A ("Secondary Phenyl Amines") with certain cyclic sulfates. Cyclic sulfates useful in the preparation
35 of "(R)-Chiral Halogenated 1-Substitutedamino-2-propanols" compounds of Formulas I-HP, I-HPC, and I-CP have a halogenated or haloalkoxy carbon adjacent to the cyclic sulfate. Some cyclic sulfates useful for the preparation of

“(R)-Chiral Halogenated 1-Substitutedamino-2-propanols” compounds of Formulas I-HP, I-HPC, and I-CP have been described by K. P. M. Vanhessche and K. B. Sharpless in Chem. Eur. J, 1997, Vol. 3, No. 4, pages 517-522 and references cited therein. (2R)-(+)-3,3,3-Trifluoro-1,2-

5 propanediol can be prepared as described in the reference cited immediately above from 3,3,3-trifluoropropene followed by separation from the predominating (2S)-(-)-3,3,3-trifluoro-1,2-propanediol. Alternatively, (2R)-(+)-3,3,3-Trifluoro-1,2-propanediol can be prepared by hydrolysis of (2R)-(+)-3,3,3-Trifluoro-2,3-epoxypropane analogous to the procedure described

10 by described by McBee and Burton in J. Am. Chem. Soc., 1952, Vol. 74, page 3022. (2R)-(+)-3,3,3-Trifluoro-1,2-propanediol is converted by reaction with a slight excess of sulfuryl chloride in the presence of 2.5 molar equivalents of imidazole, methylene chloride solvent, and at a temperature of -20 °C to give the desired (4R)-(+)-4-trifluoromethyl-2,2-dioxo-1,3,2-

15 dioxathiolane. Reaction of other (R)-Chiral haloalkyl or haloalkoxyalkyl substituted 1,2-ethanediols can afford the corresponding (4R)-substituted-2,2-dioxo-1,3,2-dioxathiolanes. Reaction of (4R)-(+)-4-trifluoromethyl-2,2-1,3,2-dioxathiolane or another (4R)-substituted-2,2-dioxo-1,3,2-dioxathiolane with a secondary amine of Formula XIII-A-H (“Secondary Heteroaryl Amines”) and Formula XIII-A (“Secondary Phenyl Amines”) in an anhydrous polar,

20 non-protic solvent such as tetrahydrofuran or acetonitrile at 25-60 °C until the reaction is complete can afford the mono-sulfate ester of a compound of Formulas I-HP, I-HPC, and I-CP. Removal of the solvent followed by addition of diethyl ether and excess 20% aqueous sulfuric acid can lead to a

25 precipitant of the crude mono-sulfate ester of a compound of Formulas I-HP, I-HPC, and I-CP. This precipitant can be filtered, the solid can be washed with ether, it can be resuspended in aqueous 20% sulfuric acid, and can be heated to 80-95 °C to give an aqueous solution of the sulfate salt of crude a compound of Formulas I-HP, I-HPC, and I-CP. Neutralization of the aqueous solution,

30 extraction with a water immiscible solvent such as diethyl ether or methylene chloride, drying the organic solvent over anhydrous magnesium sulfate, and removal of solvent can afford a compound of Formulas I-HP, I-HPC, and I-CP. Compounds of Formulas I-HP, I-HPC, and I-CP can be purified as described previously. By using a wide variety of (R)-Chiral diols, secondary

35 amines of Formula XIII-A-H (“Secondary Heteroaryl Amines”) and Formula XIII-A (“Secondary Phenyl Amines”), and reaction conditions described

herein, a large variety of compounds of Formulas I-HP, I-HPC, and I-CP may be preparable.

A particularly useful procedures to prepare Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds of the present invention in which the heteroaryl group is directly bonded is disclosed in Schemes 11 and 12. An halogenated, hydroxy containing primary alkylamine XVL ("Generic Substituted Alkylamine") formed according to Scheme 9 is itself converted by reaction with LXXI-AH ("Heteroaryl Halide") to afford secondary amine VLXX-H ("Heteroaryl Secondary Amine) using procedures disclosed in Scheme 11 and above. VLXX-H is converted to Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by alkylation chemistry with an aralkyl bromide or aralkyloxyalkyl bromide using either of two procedures disclosed in Scheme 12. Isolation and purification is effected as disclosed previously.

Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds can themselves serve as intermediates for conversion to additional compounds of this invention. Compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC and others of the present invention useful as intermediates include those in which the R₇ position substituent in Formulas I-H, I-HP, I-C, I-CP, and I-HPC is a bromo group, hydroxyl group, sulfhydryl group, bromomethyl or other bromoalkyl groups,

nitro group, amino group, methoxycarbonyl or other alkoxy carbonyl groups, cyano group, or acyl group. Other preferred compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC and the present invention useful as intermediates include those in which the R_{10} position substituent is a bromo group, hydroxyl group, sulfhydryl group, bromomethyl or other bromoalkyl groups, nitro group, amino group, methoxy carbonyl or other alkoxy carbonyl groups, cyano group, or acyl groups. Other compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC and the present invention useful as intermediates include those in which one or more of R_6 , R_7 , R_{11} , and R_{12} substituents in Formula VII is a bromo group, hydroxyl group, sulfhydryl group, bromomethyl or other bromoalkyl groups, nitro group, amino group, methoxy carbonyl or other alkoxy carbonyl groups, cyano group, or acyl groups.

A 3-bromo substituent at the R_7 position in Formula I-CP ("Polycyclic 3-Bromophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds can be reacted with a phenol to afford, as described in **Examples**, 3-phenoxy compounds of the present invention of Formula I-CP ("Polycyclic 3-Phenoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

A 3-bromo substituent at the R_7 position in Formula I-HP and I-HPC ("Polycyclic 3-Bromophenyl and 3-Bromoheteroaryl/Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") can, as shown in Scheme 4, be reacted with a phenol to afford, as described in **Examples**, additional compounds of the present invention of Formula I-HP and I-HPC ("Polycyclic 3-Aryloxyaryl, 3-Heteroaryloxyaryl, 3-Heteroaryloxyheteroaryl, 3-Aryloxyheteroaryl, 3-Arylthioaryl, 3-Heteroarylthioaryl, 3-Heteroarylthioheteroaryl, and 3-Arylthioheteroaryl Aryl and Heteroaryl/Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

A 3-bromo substituent at the R_7 position in Formula I-CP ("Polycyclic 3-Bromophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") can be reacted, as shown in Scheme 7, with an aryl borinate or an aryl tin to afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Arylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Scheme 8 discloses the conversion of a 3-bromo substituent at the R_7 position in Formula I-CP ("Polycyclic 3-Bromophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with a primary or

secondary amine to afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3- R₂₂aminophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-bromo substituent at the R₁₀ position in Formula I-CP ("Polycyclic 3-Bromophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an aryl borinate can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Arylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-bromo substituent at the R₁₀ position in Formula I-CP ("Polycyclic 3-Bromophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with a heteroaryl dibutyl tin compound can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Heteroarylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-bromomethyl substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Bromomethylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") by reaction with an aryl borinate can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Arylmethylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Scheme 5 discloses the conversion of a 3-hydroxyl substituent at the R₇ position in Formula I-HP and I-HPC ("Polycyclic 3-Hydroxyphenyl and 3-Hydroxyheteroaryl/Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") by reaction with an aryl bromide or heteroaryl bromide to afford, as described in **Examples**, additional compounds of the present invention of Formula I-HP and I-HPC ("Polycyclic 3-Aryloxyaryl, 3-Heteroaryloxyaryl, 3-Heteroaryloxyheteroaryl, and 3-Aryloxyheteroaryl Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-hydroxyl substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Hydroxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an aryl bromide can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Phenoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-hydroxyl substituent at the R₇ position in Formula I-HP and I-HPC ("Polycyclic 3-Hydroxyphenyl and 3-Hydroxyheteroaryl/Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols")

compounds by reaction with an aralkyl bromide or heteroaralkyl bromide can afford, as described above for Scheme 5 and in **Examples**, additional compounds of the present invention of Formula I-HP and I-HPC ("Polycyclic 3-Aralkyloxyaryl, 3-Heteroaralkyloxyaryl, 3-Heteroaralkyloxyheteroaryl, and 3-Aralkyloxyheteroaryl Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-hydroxyl substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Hydroxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an aralkyl bromide can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Aralkyloxyaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-hydroxyl substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Hydroxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an R₁₋₇-bromide can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3- R₁₋₇-oxyaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-thio substituent at the R₇ position in Formula I-CP ("Polycyclic 3-thiophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an R₁₋₇-bromide can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3- R₁₋₇-thiaaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"). "Polycyclic 3- R₁₋₇-thiaaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols" can be oxidized to sulfonyl compounds of Formula I-CP ("Polycyclic 3- R₁₋₇-sulfonylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-nitro substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Nitrophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by hydrogenation can afford, as described in

Examples. additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Aminophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"). "Polycyclic 3-Aminophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols" can be acylated to acyl amide compounds of
5 Formula I-CP ("Polycyclic 3-R₁₇-C(O)amidophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-amino substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Aminophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with carbonyl compounds can afford, as
10 described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-(Saturated Nitrogen Heterocycl-1yl)aryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols" and ("Polycyclic 3-(Unsaturated Nitrogen Heterocycl-1yl)aryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

15 Conversion of a 3-methoxycarbonyl substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Carbomethoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with amination reagents can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Carboxamidophenyl (R)-
20 Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-cyano substituent at the R₇ position in Formula I-CP ("Polycyclic 3-Cyanophenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with organometallic reagents can afford, as described in **Examples**, additional compounds of the present invention of
25 Formula I-CP ("Polycyclic 3-Acylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"). Said "Polycyclic 3-Acylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols" can be reduced to hydroxyl compounds of Formula I-CP ("Polycyclic 3-hydroxysubstitutedmethylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

30 Conversion of a 3-methoxycarbonyl substituent at the R₁₀ position in Formula I-CP ("Polycyclic 3-Carbomethoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with amination reagents can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP "Polycyclic 3-Carboxamdophenyl (R)-
35 Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-methoxycarbonyl substituent at the R_{10} position in Formula I-CP ("Polycyclic 3-Carbomethoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an organometallic reagent can afford, as described in **Examples**, additional
5 compounds of the present invention of Formula I-CP "Polycyclic 3-(bis- R_{20} -hydroxymethyl)aryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-methoxycarbonyl substituent at the R_{10} position in Formula I-CP ("Polycyclic 3-Carbomethoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with lithium
10 aluminum hydride can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-Hydroxymethylphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Conversion of a 3-methoxycarbonyl substituent at the R_{10} position in Formula I-CP ("Polycyclic 3-Carbomethoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction with an alkylation
15 reagent can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-(bis- R_{21} -hydroxymethyl)phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").
20

Conversion of a 3-methoxycarbonyl substituent at the R_{10} position in Formula I-CP ("Polycyclic 3-Carbomethoxyphenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") compounds by reaction initially with an
25 amidation reagent and then an R_{20} -organometallic reagent can afford, as described in **Examples**, additional compounds of the present invention of Formula I-CP ("Polycyclic 3-(R_{20} -carbonyl)phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols").

Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and
30

Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") and other compounds of this invention possessing hydroxyl, thiol, and amine functional groups can be converted to a wide variety derivatives. The hydroxyl group, wherein R_{16} is a hydrogen and

5 X is oxy, of compounds of Formulas I-H, I-HP, I-HPC, I-C, and I-CP can be readily converted to esters of carboxylic, sulfonic, carbamic, phosphonic, and phosphoric acids. Acylation to form a carboxylic acid ester is readily effected using a suitable acylating reagent such as an aliphatic acid anhydride or acid chloride. The corresponding aryl and heteroaryl acid anhydrides and acid

10 chlorides can also be used. Such reactions are generally carried out using an amine catalyst such as pyridine in an inert solvent. In like manner, compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP that have at least one hydroxyl group present in the form of an alcohol or phenol can be acylated to its corresponding esters. Similarly, carbamic acid

15 esters (urethans) can be obtained by reacting any hydroxyl group with isocyanates and carbamoyl chlorides. Sulfonate, phosphonate, and phosphate esters can be prepared using the corresponding acid chloride and similar reagents. Compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP that have at least one thiol group present can be

20 converted to the corresponding thioesters derivatives analogous to those of alcohols and phenols using the same reagents and comparable reaction conditions. Compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP that have at least one primary or secondary amine group present can be converted to the corresponding amide derivatives.

25 Amides of carboxylic acids can be prepared using the appropriate acid chloride or anhydrides with reaction conditions analogous to those used with alcohols and phenols. Ureas of the corresponding primary or secondary amine can be prepared using isocyanates directly and carbamoyl chlorides in the presence of an acid scavenger such as triethylamine or pyridine. Sulfonamides can be

30 prepared from the corresponding sulfonyl chloride in the presence of aqueous sodium hydroxide. Suitable procedures and methods for preparing these derivatives can be found in House's Modern Synthetic Reactions, W. A. Benjamin, Inc., Shriner, Fuson, and Curtin in The Systematic Identification of Organic Compounds, 5th Edition, John Wiley & Sons, and Fieser and

35 Fieser in Reagents for Organic Synthesis, Volume 1, John Wiley & Sons. Reagents of a wide variety that can be used to derivatize hydroxyl, thiol, and

amines of compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP are available from commercial sources or the references cited above, which are incorporated herein by reference.

- Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") and other compounds of this invention possessing hydroxyl, thiol, and amine functional groups can be alkylated to a wide variety derivatives. The hydroxyl group, wherein R_{16} is a hydrogen and X is oxy, of compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP can be readily converted to ethers. Alkylation to form an ether is readily effected using a suitable alkylating reagent such as an alkyl bromide, alkyl iodide or alkyl sulfonate. The corresponding aralkyl, heteroaralkyl, alkoxyalkyl, aralkyloxyalkyl, and heteroaralkyloxyalkyl bromides, iodides, and sulfonates can also be used. Such reactions are generally carried out using an alkoxide forming reagent such as sodium hydride, potassium t-butoxide, sodium amide, lithium amide, and n-butyl lithium using an inert polar solvent such as DMF, DMSO, THF, and similar, comparable solvents. In like manner, compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP that have at least one hydroxyl group present in the form of an alcohol or phenol can be alkylated to their corresponding ethers. Compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP that have at least one thiol group present can be converted to the corresponding thioether derivatives analogous to those of alcohols and phenols using the same reagents and comparable reaction conditions. Compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP that have at least one primary, secondary or tertiary amine group present can be converted to the corresponding quaternary ammonium derivatives. Quaternary ammonium derivatives can be prepared using the appropriate bromides, iodides, and sulfonates analogous to those used with alcohols and phenols. Conditions involve reaction of the amine by warming it

with the alkylating reagent with a stoichiometric amount of the amine (i.e., one equivalent with a tertiary amine, two with a secondary, and three with a primary). With primary and secondary amines, two and one equivalents, respectively, of an acid scavenger are used concurrently. Tertiary amines can be prepared from the corresponding primary or secondary amine by reductive alkylation with aldehydes and ketones using reduction methods 1, 2, or 3 as shown in Scheme 1. Suitable procedures and methods for preparing these derivatives can be found in House's Modern Synthetic Reactions, W. A. Benjamin, Inc., Shriner, Fuson, and Curtin in The Systematic Identification of Organic Compounds, 5th Edition, John Wiley & Sons, and Fieser and Fieser in Reagents for Organic Synthesis, Volume 1, John Wiley & Sons. Perfluoroalkyl derivatives can be prepared as described by DesMarteau in J. Chem. Soc. Chem. Commun. 2241 (1998). Reagents of a wide variety that can be used to derivatize hydroxyl, thiol, and amines of compounds of Formulas I-H, I-HP, I-C, I-CP, I-HPC, Cyclo I-H, Cyclo I-C, and Cyclo I-CP are available from commercial sources or the references cited above, which are incorporated herein by reference.

Formula I-H ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-alkanols"), Formula I-HP ("Generic Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-propanols"), Formula I-HPC ("Polycyclic Aryl-Heteroaryl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols"), Formula I-C ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-(n+1)-Alkanols"), and Formula I-CP ("Polycyclic Phenyl (R)-Chiral Halogenated 1-Substitutedamino-2-Propanols") and certain other compounds of this invention can be converted, according to Scheme 6, to the corresponding cyclic derivatives represented by "Tricyclic *tertiary*-oxyalkylamines" and exemplified by Formulas Cyclo I-H ("Polycyclic Aryl and Heteroaryl (R)-Chiral Halogenated (N+1)-Cycloazaalkoxy"), Cyclo I-C ("Polycyclic Aryl Phenyl (R)-Chiral Halogenated (N+1)-Cycloazaalkoxy") and Cyclo I-CP ("Polycyclic Phenyl Phenyl (R)-Chiral Halogenated Cycloazaalkoxy"). The hydroxyl group, wherein R₁₆ is a hydrogen and X is oxy, of compounds of Formulas I-H, I-HP, I-C, I-CP, and I-HPC can be cyclized to corresponding cyclic ethers. Compounds suitable for cyclization will normally have at least one leaving group within 5 to 10 continuous atoms of the hydroxyl group wherein R₁₆ is a hydrogen and X is oxy. Most preferably the leaving group will be

within 5 to 7 atoms of the hydroxyl group so as to form a 6 to 8 membered ring heteroatom containing ring. When the leaving group is part of an aromatic ring system, the leaving group will be preferably in an ortho position.

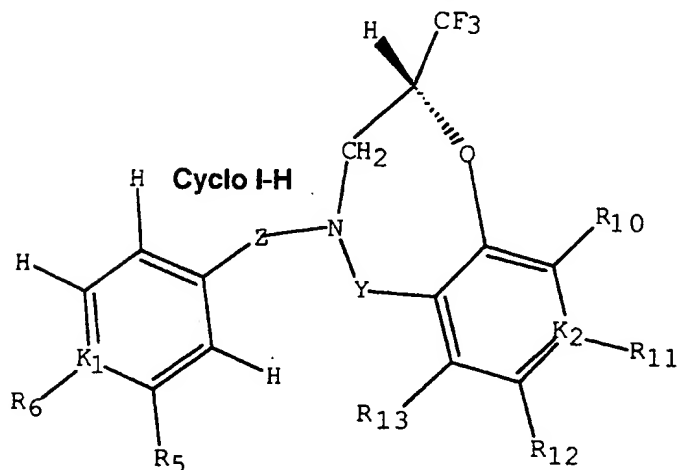
Suitable leaving groups generally include halides, sulfates, sulfonates,

- 5 trisubstituted amino, disubstituted sulfonium, diazonium, and like, and, in the case of aromatic systems, also includes nitro, alkoxy, aryloxy, heteroaryloxy, and alkylthio.

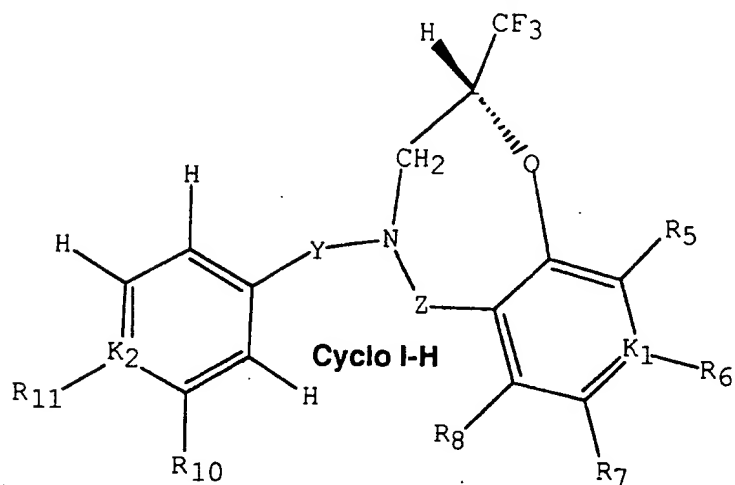
- The cyclization reaction to form "Tricyclic *tertiary*-oxyalkylamines" of Formulas Cyclo I-H, Cyclo I-C and Cyclo I-CP can be accomplished by aromatic and aliphatic nucleophilic substitution reactions such as those disclosed in March's Advanced Organic Chemistry, 4th Edition, John Wiley & Sons, especially at pages 293-412 and 649-658 and the references cited therein, which are incorporated herein by reference. Hydroxyl containing suitably substituted compounds can be converted to a cyclic analog by heating a suitably substituted compound under anhydrous conditions in a suitable solvent, such as dimethylformamide, dimethylsulfoxide, N-methylpyrrolidone, tetraglyme, or hexamethylphosphoramide, in the presence of a suitable base such as potassium carbonate, cesium carbonate, sodium hydroxide, potassium *tertiary*-butoxide, or lithium diisopropylamide. Alternately, sodium amide in anhydrous ammonia solvent can be used. Temperatures in the range of -20 °C to 200 °C can be used for time periods of 30 minutes to more than 24 hours.

- The preferred temperature can be selected by standard synthetic chemical technique balancing maximum yield, maximum purity, cost, ease of isolation and operation, and time required. Isolation of the "Tricyclic *tertiary*-oxyalkylamines" can be effected as described above for other *tertiary*-oxyalkylamines. Representative "Tricyclic *tertiary*-oxyalkylamines" prepared using the methodology described above are included in Table 5.

- The following examples are provided to illustrate the present invention and are not intended to limit the scope thereof. Those skilled in the art will readily understand that known variations of the conditions and processes of the following preparative procedures can be used to prepare these compounds.

Table 5. Structure of Substituted Tricyclic^{tertiary}-2-oxyalkylamines.

| <u>Y</u> | <u>Z</u> | <u>R₅</u> | <u>K₁-R₆</u> | <u>R₁₀</u> | <u>K₂-R₁₁</u> | <u>R₁₂</u> | <u>R₁₃</u> |
|---------------|----------|-------------------------|------------------------------------|-----------------------|-------------------------------------|-----------------------|-----------------------|
| CH_2 | - | 4-chloro-3-ethylphenoxy | C-H | H | C- CF_3 | H | H |
| CH_2 | - | 4-chloro-3-ethylphenoxy | N | H | C- CF_3 | H | H |
| CH_2 | - | 4-chloro-3-ethylphenoxy | C-H | H | C- H | CF_3 | H |
| CH_2 | - | 4-chloro-3-ethylphenoxy | N | H | C- H | CF_3 | H |
| CH_2 | - | 4-chloro-3-ethylphenoxy | C-H | H | N | CF_3 | H |
| - | - | 4-chloro-3-ethylphenoxy | C-H | H | C- CF_3 | H | H |
| - | - | 4-chloro-3-ethylphenoxy | N | H | C- CF_3 | H | H |
| - | - | 4-chloro-3-ethylphenoxy | C-H | H | C- H | CF_3 | H |
| - | - | 4-chloro-3-ethylphenoxy | N | H | C- H | CF_3 | H |
| - | - | 4-chloro-3-ethylphenoxy | C-H | H | N | CF_3 | H |

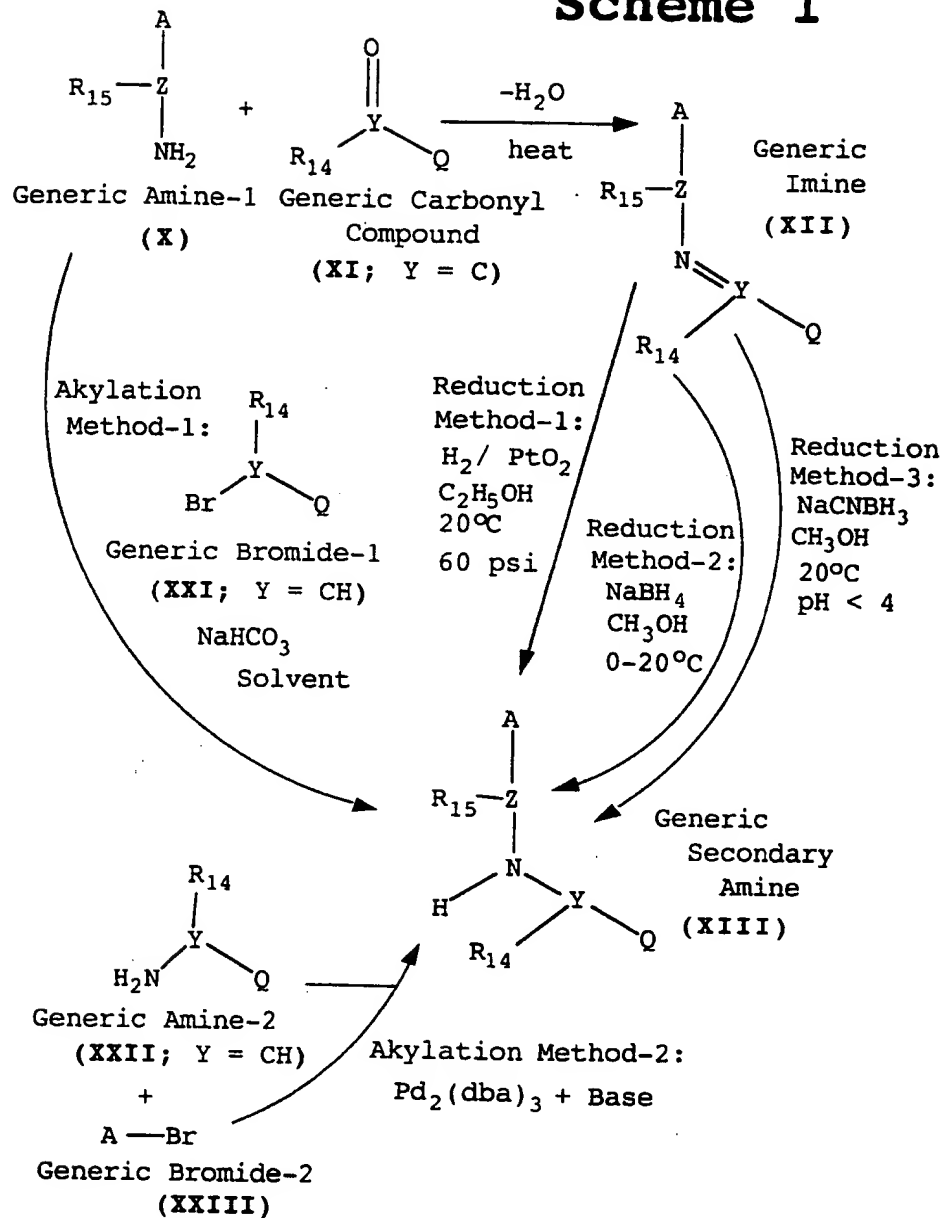
Table 5. (cont.) Structure of Substituted Tricyclic *tertiary*-2-oxyalkylamines.

| <u>Y</u> | <u>Z</u> | <u>R₅</u> | <u>K₁-R₆</u> | <u>R₁₀</u> | <u>K₂-R₁₁</u> | <u>R₇</u> | <u>R₈</u> |
|-----------------|----------|-------------------------|------------------------------------|------------------------------------|-------------------------------------|----------------------|----------------------|
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | OCF ₂ CF ₂ H | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | N | OCF ₂ CF ₂ H | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | OCF ₂ CF ₂ H | N | H | H |
| CH ₂ | - | phenoxy | C-H | OCF ₂ CF ₂ H | C-H | H | H |
| CH ₂ | - | phenoxy | N | OCF ₂ CF ₂ H | C-H | H | H |
| CH ₂ | - | phenoxy | C-H | OCF ₂ CF ₂ H | N | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | CF ₂ CF ₃ | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | N | CF ₂ CF ₃ | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | CF ₂ CF ₃ | N | H | H |
| CH ₂ | - | phenoxy | C-H | CF ₂ CF ₃ | C-H | H | H |
| CH ₂ | - | phenoxy | N | CF ₂ CF ₃ | C-H | H | H |
| CH ₂ | - | phenoxy | C-H | CF ₂ CF ₃ | N | H | H |

Table 5. (cont.) Structure of Substituted Tricyclic *tertiary*-2-oxyalkylamines.

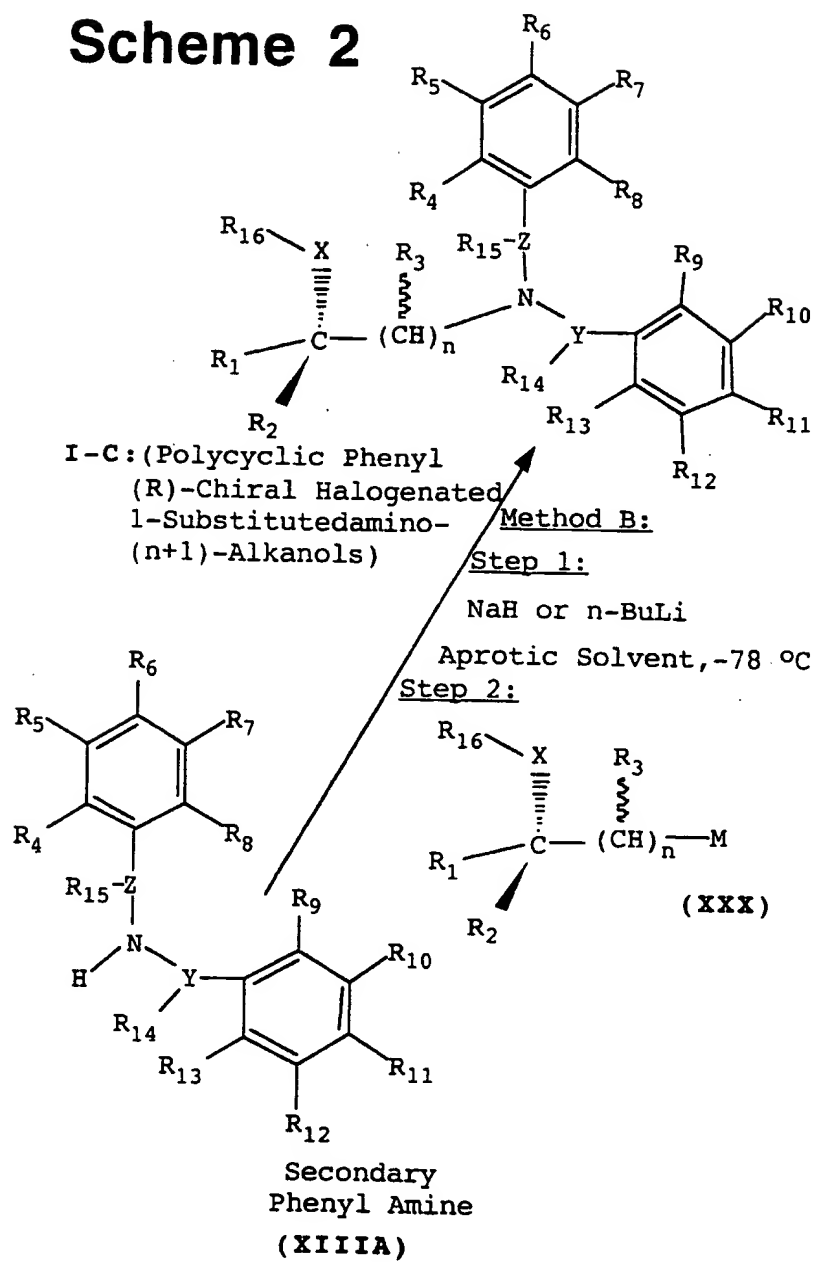
| <u>Y</u> | <u>Z</u> | <u>R₅</u> | <u>K₁-R₆</u> | <u>R₁₀</u> | <u>K₂-R₁₁</u> | <u>R₇</u> | <u>R₈</u> |
|-----------------|----------|-------------------------|------------------------------------|------------------------------------|-------------------------------------|----------------------|----------------------|
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | CF ₃ | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | N | CF ₃ | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | CF ₃ | N | H | H |
| CH ₂ | - | phenoxy | C-H | CF ₃ | C-H | H | H |
| CH ₂ | - | phenoxy | N | CF ₃ | C-H | H | H |
| CH ₂ | - | phenoxy | C-H | CF ₃ | N | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | OCF ₂ CF ₂ H | C-H | H | F |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | N | OCF ₂ CF ₂ H | C-H | H | F |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | OCF ₂ CF ₂ H | N | H | F |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | 2-furyl | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | N | 2-furyl | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | 2-furyl | N | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | SCF ₃ | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | N | SCF ₃ | C-H | H | H |
| CH ₂ | - | 4-chloro-3-ethylphenoxy | C-H | SCF ₃ | N | H | H |

Scheme 1

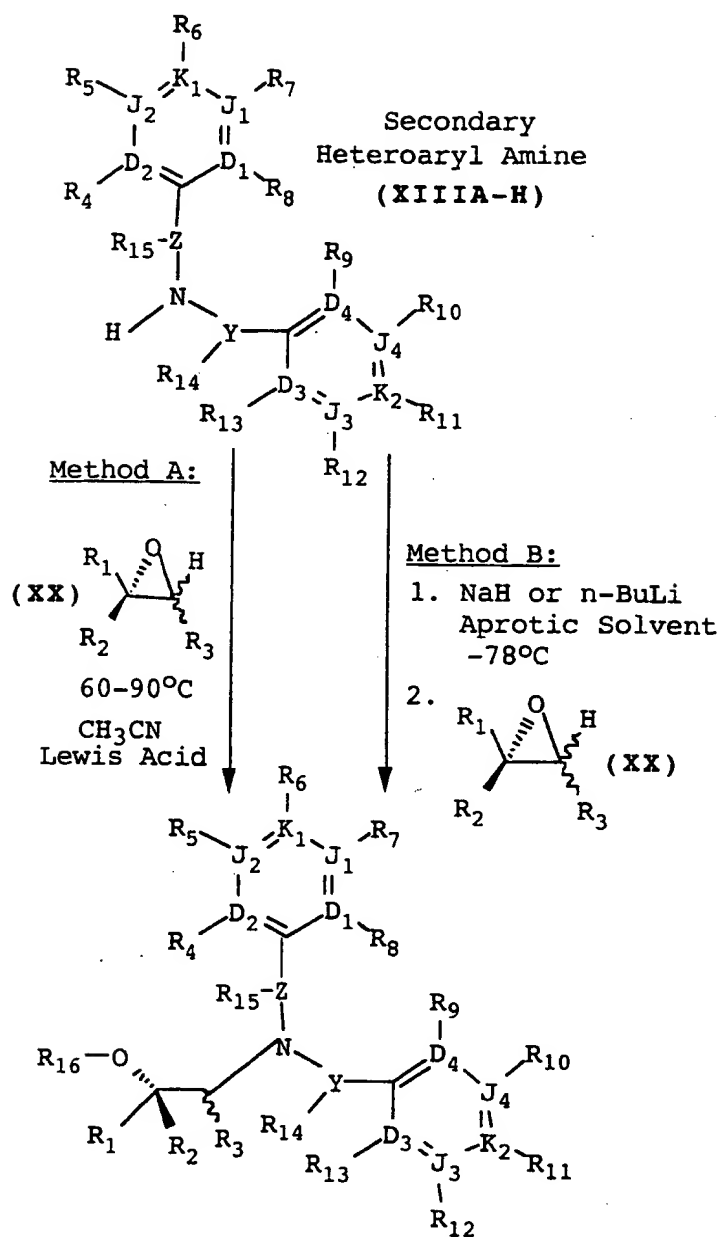


109

Scheme 2

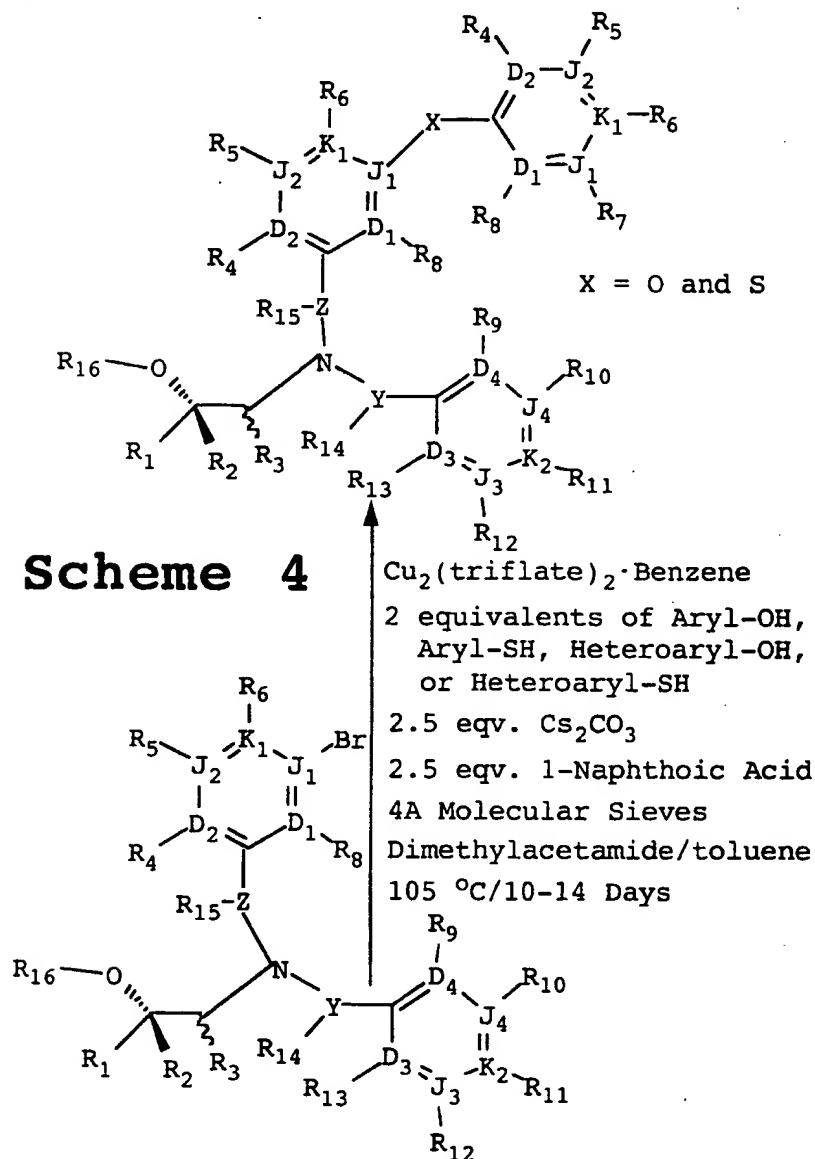


Scheme 3



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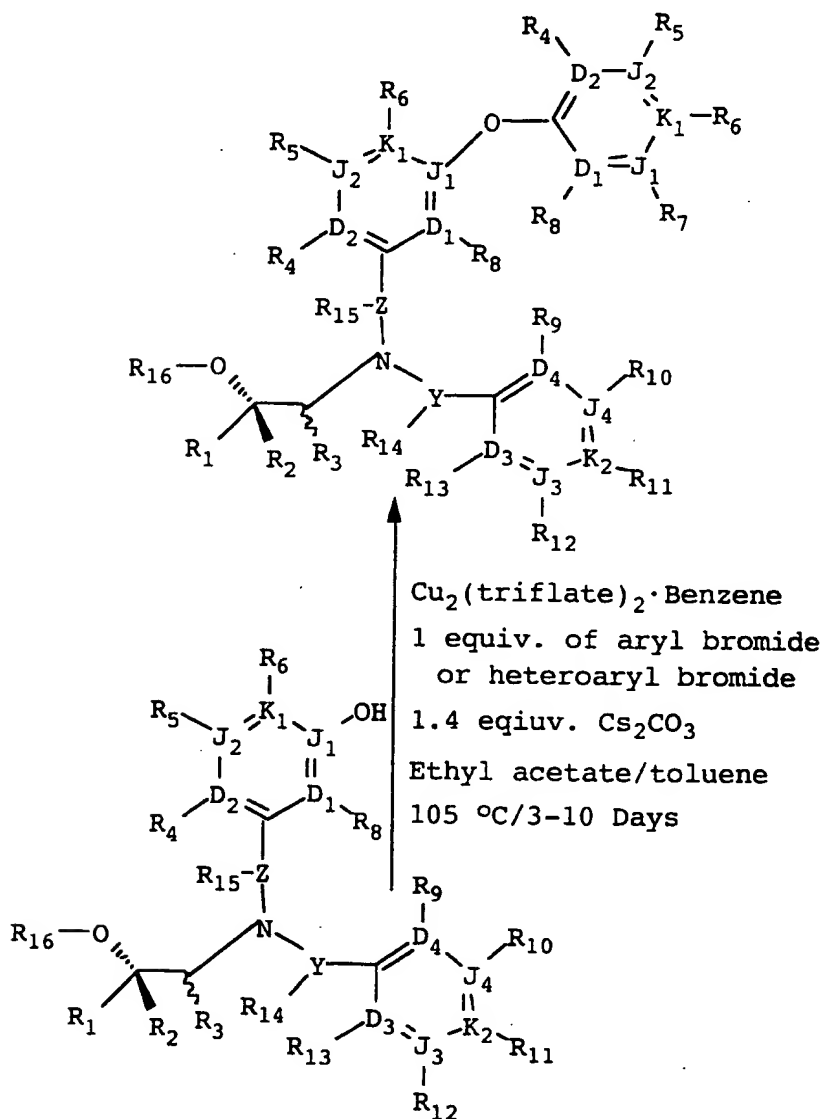
I-HP/I-HPC:(Generic Polycyclic 3-Aryloxyaryl,
3-Heteroaryloxyaryl,3-Heteroaryloxyheteroaryl,
3-Aryloxyheteroaryl, 3-Arylthioaryl,
3-Heteroarylthioaryl,3-Heteroarylthioheteroaryl,
3-Arylthioheteroaryl Aryl and Heteroaryl/Aryl-
Heteroaryl (R)-Chiral Halogenated
1-Substitutedamino-2-Propanols)



I-HP/I-HPC:(Generic Polycyclic 3-Bromo Aryl and
Heteroaryl/Aryl-Heteroaryl (R)-Chiral
Halogenated 1-Substitutedamino-2-Propanols)

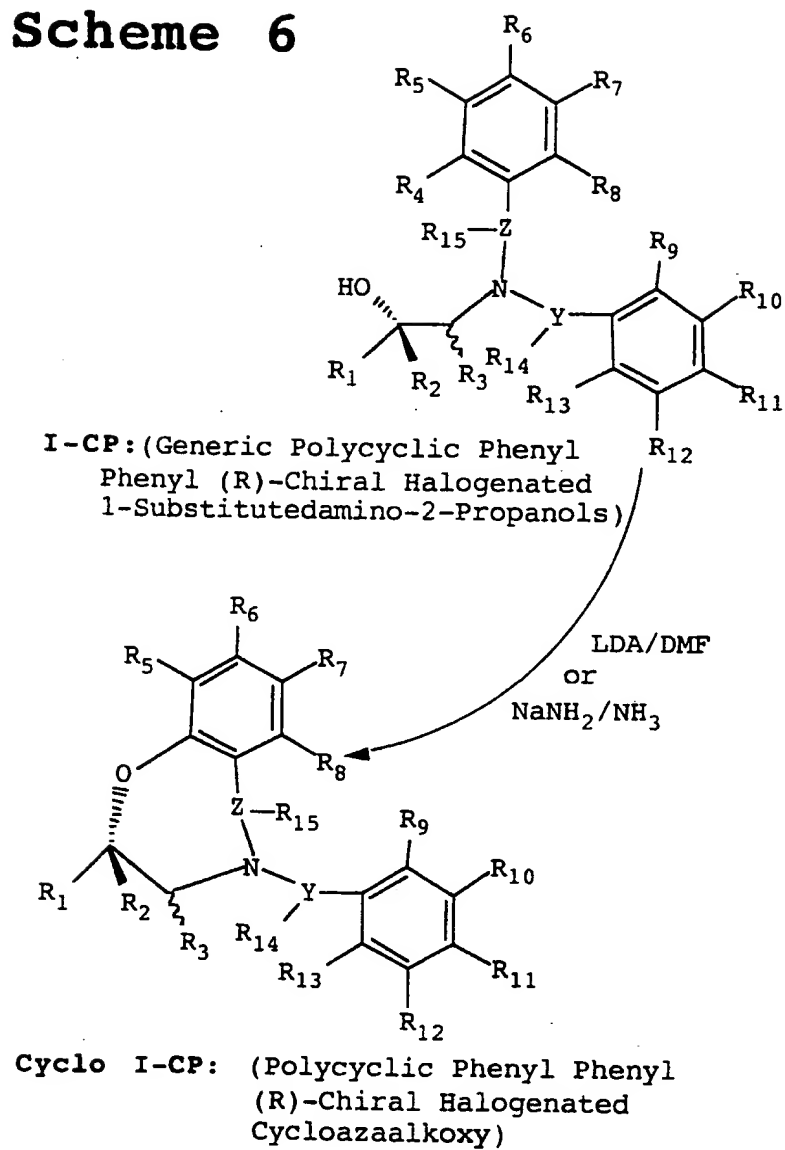
Scheme 5

I-HP/I-HPC: (Generic Polycyclic 3-Aryloxyaryl,
3-Heteroaryloxyaryl 3-Aryloxyheteroaryl
or 3-Heteroaryloxyheteroaryl
Aryl-Heteroaryl (R)-Chiral Halogenated
1-Substitutedamino-2-Propanols)



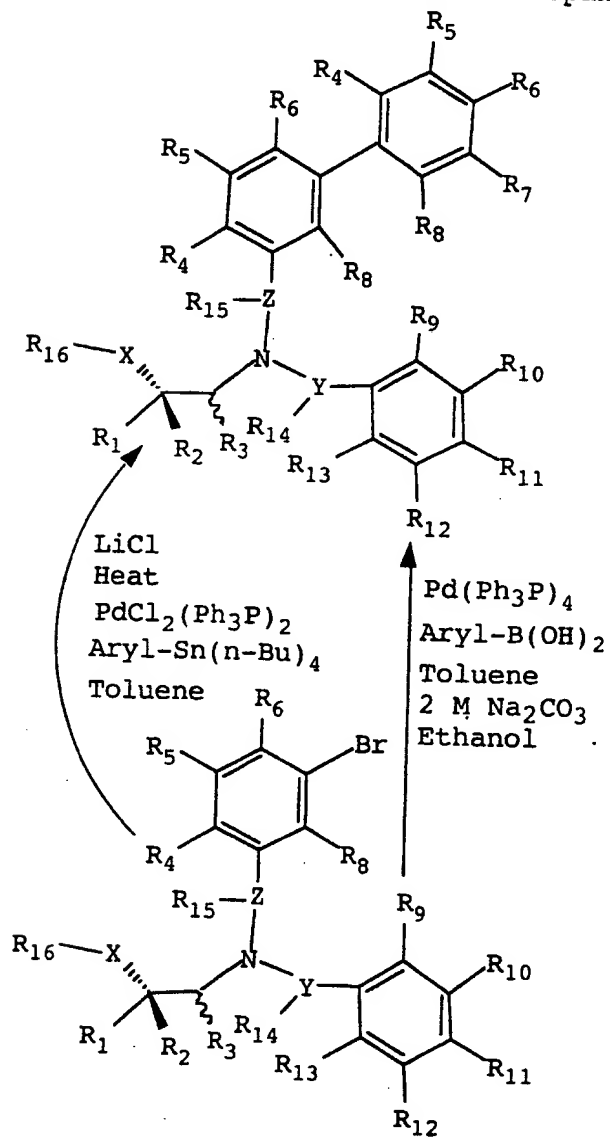
I-HP/I-HPC: (Generic Polycyclic 3-Hydroxy Aryl
and Heteroaryl/Aryl-Heteroaryl (R)-Chiral
Halogenated 1-Substitutedamino-2-Propanols)

113

Scheme 6

Scheme 7

I-CP: (Polycyclic 3-Arylphenyl
Phenyl (R)-Chiral Halogenated
1-Substitutedamino-2-Propanols)

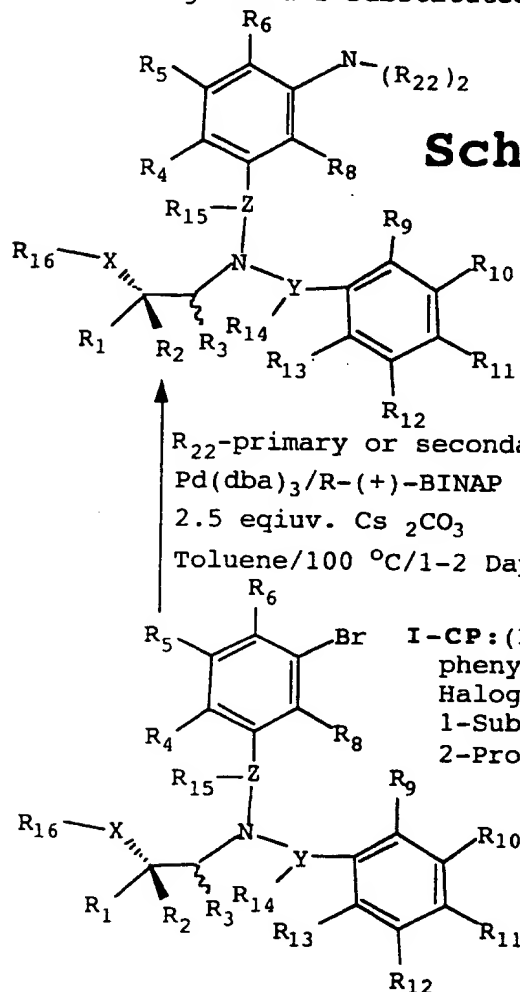


I-CP: (Polycyclic 3-Bromophenyl
Phenyl (R)-Chiral Halogenated
1-Substitutedamino-2-Propanols)

NOTE: Use of Heteroaryl-B(OH)₂ will give
the heteroarylmethyl analog of I-CP.

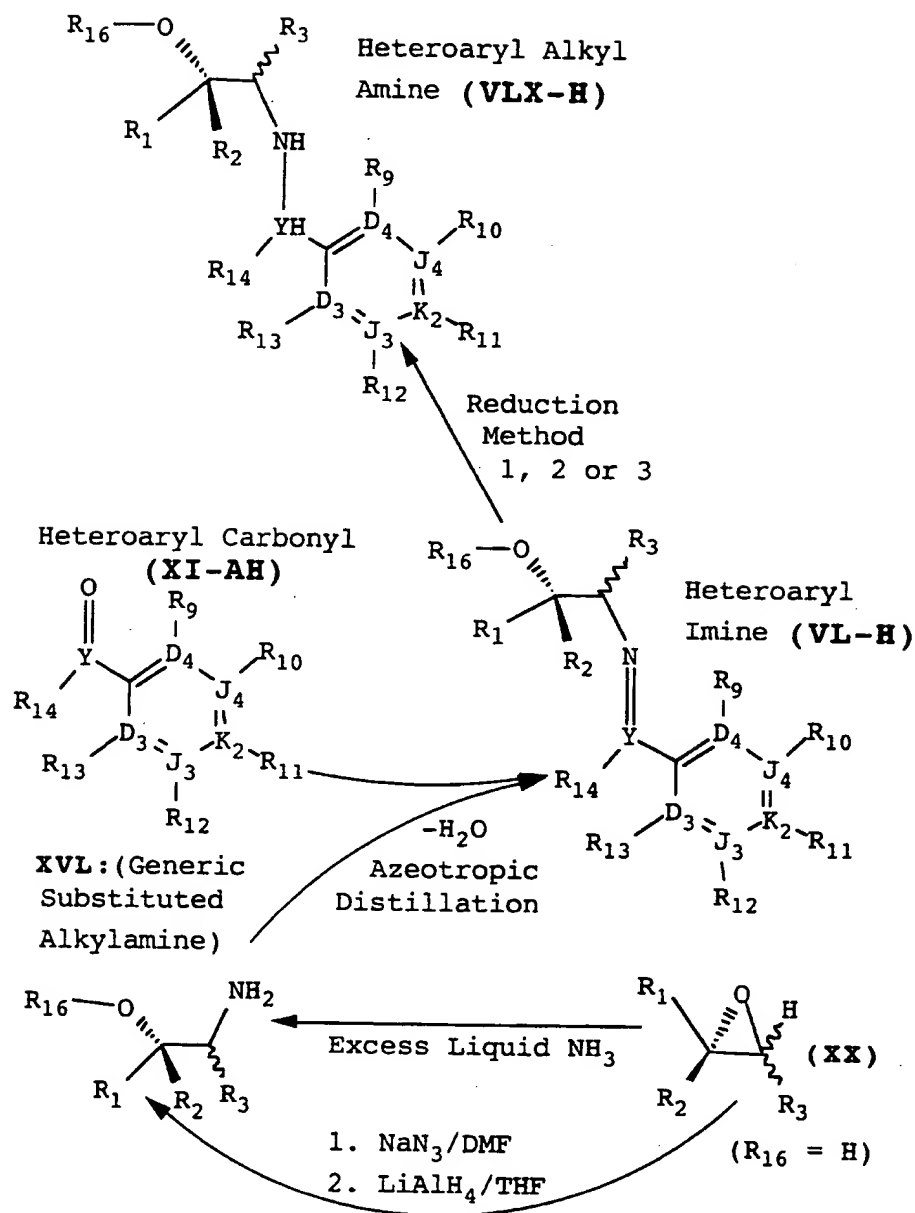
I-CP:(Polycyclic 3- R_{22} -aminophenyl Phenyl (R)-Chiral
Halogenated 1-Substitutedamino-2-Propanols)

Scheme 8

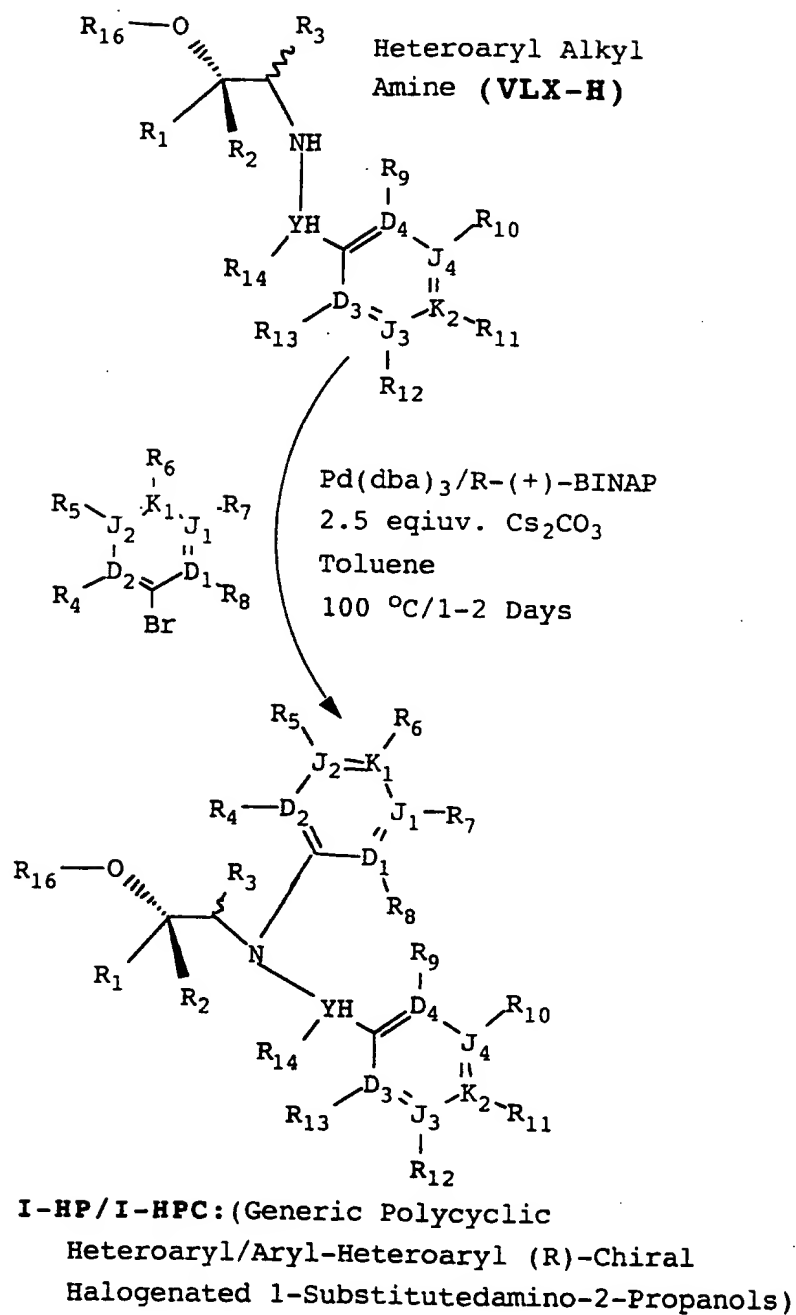


R_{22} is selected independently from any one or two of the following groups: hydrido, hydroxy, aryloxy, alkyl, alkenyl, alkynyl, aryl, aralkyl, alkoxyalkyl, aralkoxyalkyl, alkylsulfinylalkyl, haloalkyl, haloalkenyl, halocycloalkyl, halocycloalkoxy, halocycloalkoxyalkyl, arylsulfinylalkyl, arylsulfonylalkyl, alkylamino, cycloalkylsulfinylalkyl, cycloalkylsulfonylalkyl, heteroaryl sulfonylalkyl, heteroaryl sulfinylalkyl, aralkylsulfinylalkyl, aralkylsulfonylalkyl, hydroxyalkyl, amino, alkoxy, alkylsulfonylalkyl, aralkylthioalkyl, heteroaralkylthioalkyl, heteroaryloxyalkyl, alkenyloxyalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, heteroaryl, halocycloalkenyloxyalkyl, heteroarylalkyl, aryloxyalkyl, halocycloalkenyl, and heteroarylthioalkyl.

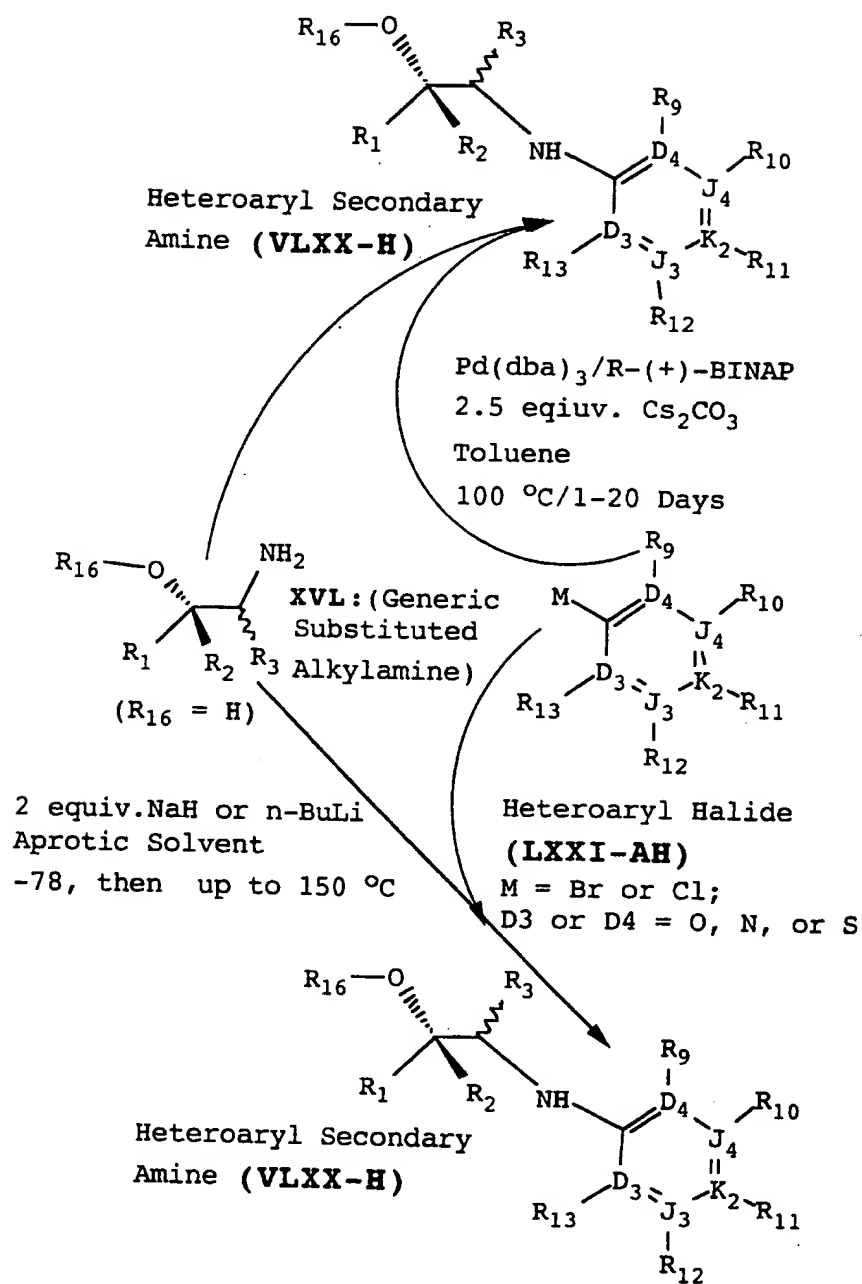
Scheme 9



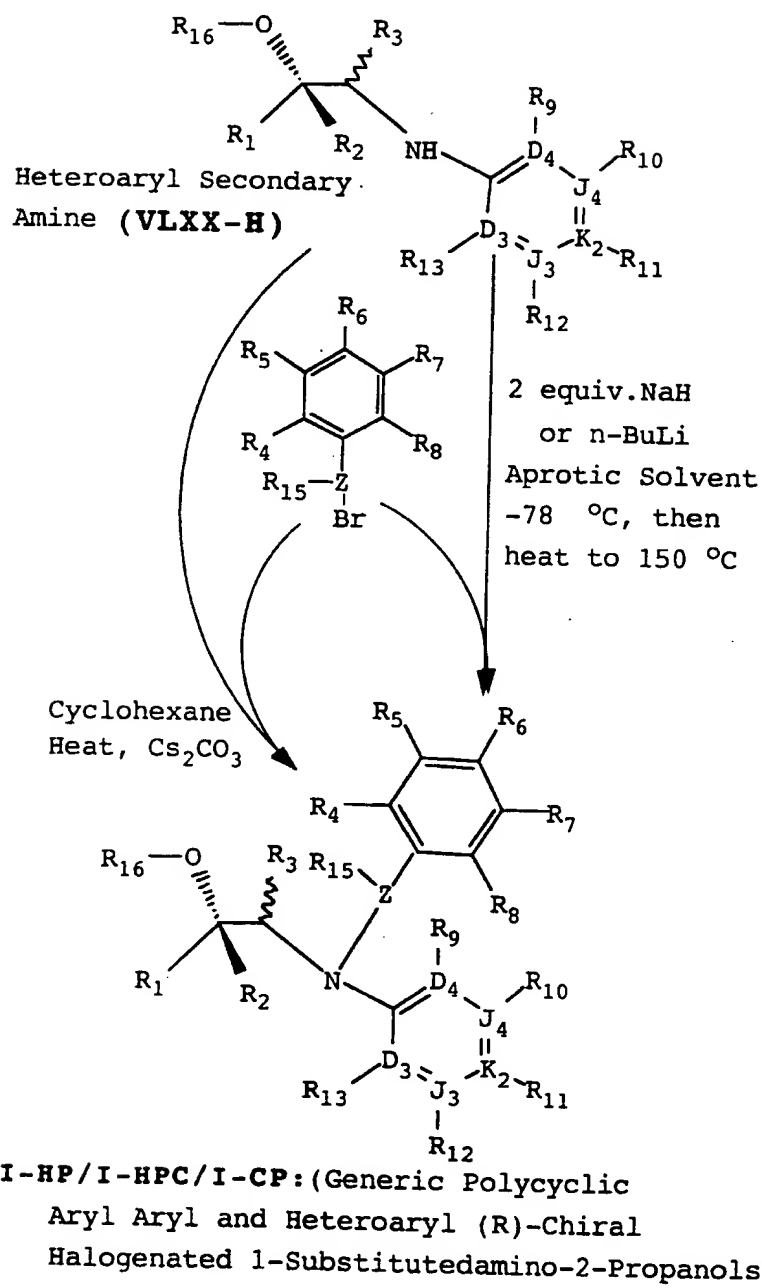
Scheme 10



Scheme 11

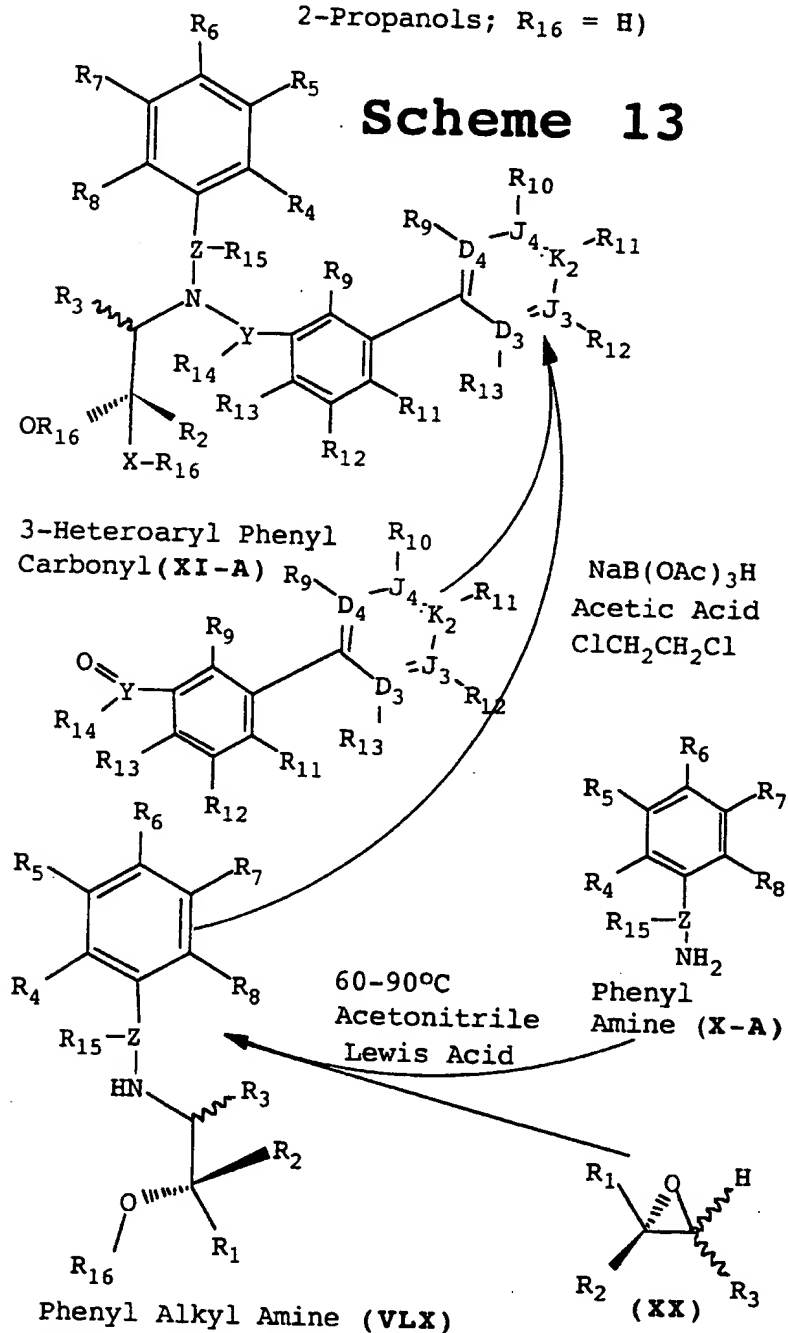


Scheme 12



I-CP: (Polycyclic 3-Heteroarylaryl Phenyl
(R)-Chiral Halogenated 1-Substitutedamino-
2-Propanols; $R_{16} = H$)

Scheme 13

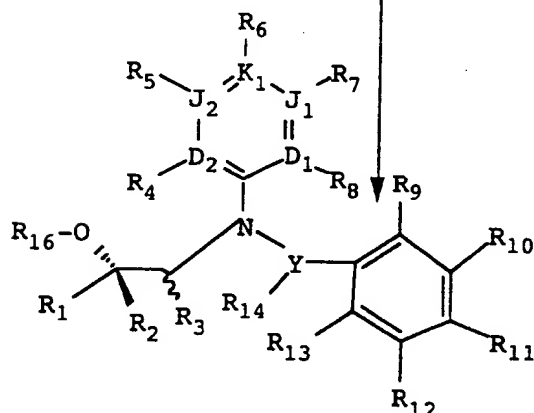
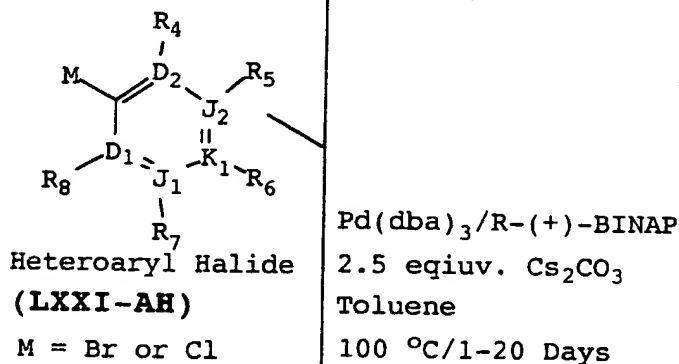
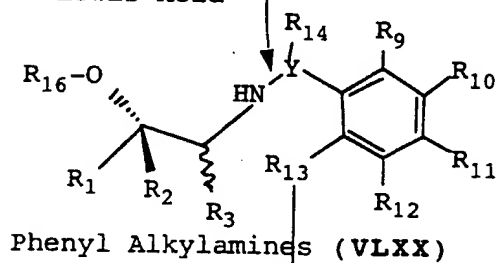
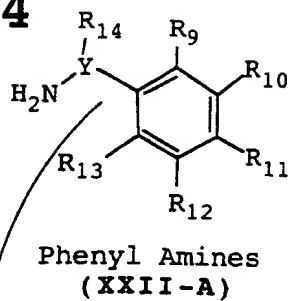
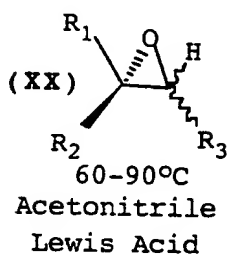


NOTE: Heteroaryl Analogs Can Be Prepared Using Heteroaryl Analogs of **X-A**, **VLX**, and **XI-A**.

121

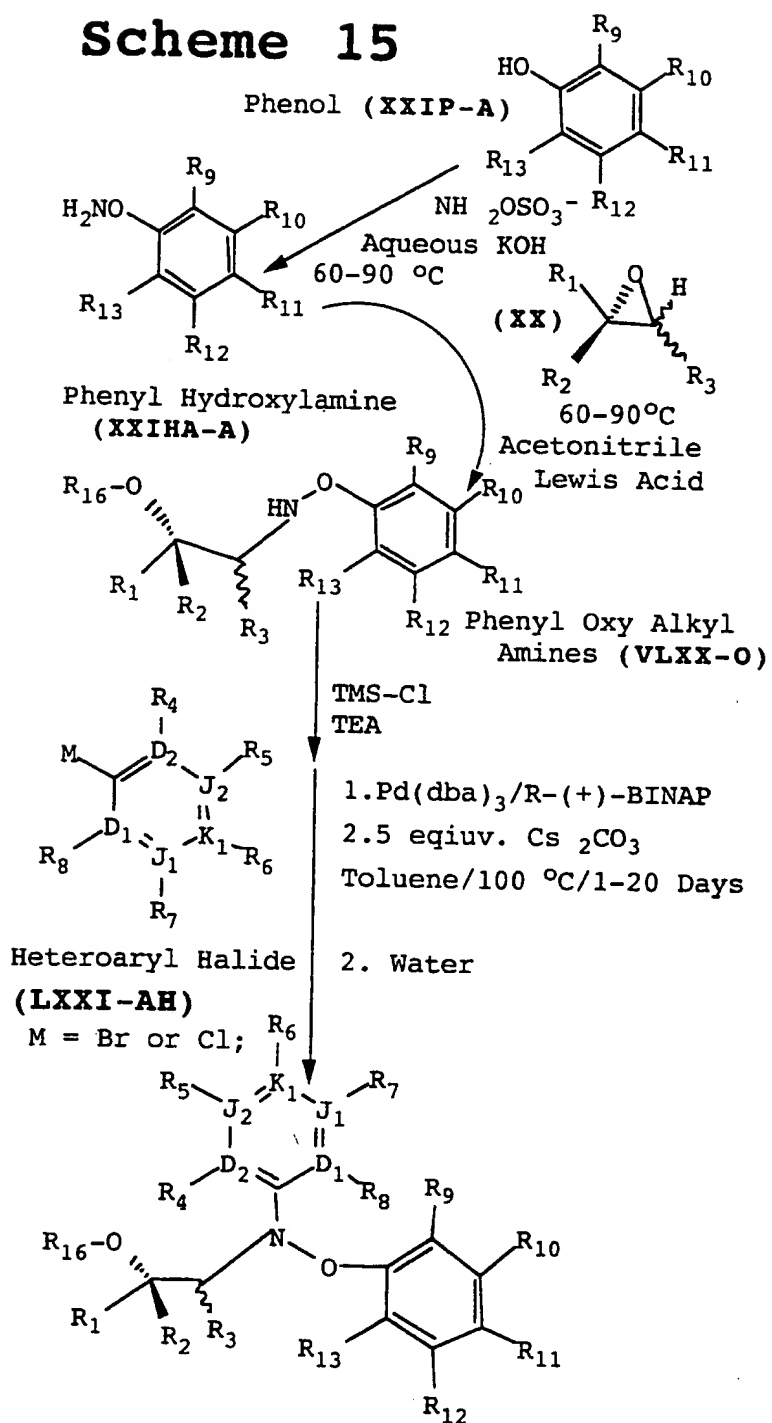
Scheme 14

Method A:



I-HPC: Polycyclic Aryl-Heteroaryl (R)-Chiral
Halogenated 1-Substitutedamino-2-Propanols
when R₁₆ equals H) NOTE: Aryl Analogs
(I-CP) of (I-HPC) Can Be Prepared by Starting
With Aryl Bromide Analogs of (LXXI-AH).

Scheme 15



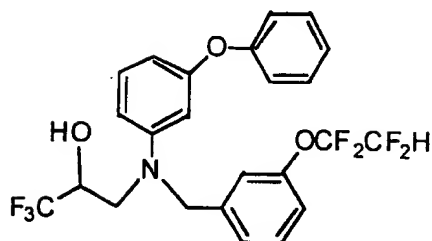
I-HPC: (Polycyclic Heteroaryl-Aryl Phenyl
(R)-Chiral Halogenated 1-Substitutedamino-2-Propanols
when R₁₆ = H and Y = O) NOTE: Diaryl (I-CP) and
Diheteroaryl (I-HP) Analogs Can Be Prepared by Using
Aryl Bromide and Heteroaryl-OH, respectively.

The following examples are provided to illustrate the present invention and are not intended to limit the scope thereof. Without further elaboration, it is believed that one skilled in the art can, using the preceding descriptions, utilize the present invention to its fullest extent. Therefore the following preferred specific embodiments are to be construed as merely illustrative and not limitative of the remainder of the disclosure in any way whatsoever. Compounds containing multiple variations of the structural modifications illustrated in the preceding schemes or the following Examples are also contemplated. Those skilled in the art will readily understand that known variations of the conditions and processes of the following preparative procedures can be used to prepare these compounds.

One skilled in the art may use these generic methods to prepare the following specific examples, which have been or may be properly characterized by ¹H NMR and mass spectrometry. These compounds also may be formed in vivo.

The following examples contain detailed descriptions of the methods of preparation of compounds of Formula V-H. These detailed descriptions fall within the scope and are presented for illustrative purposes only and are not intended as a restriction on the scope of the invention. All parts are by weight and temperatures are Degrees centigrade unless otherwise indicated.

EXAMPLE 1



5

(2*R,S*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol

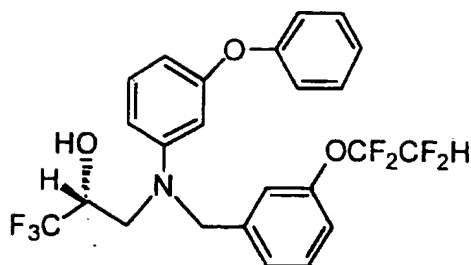
EX-1A) To a solution of 3-(1,1,2,2-tetrafluoroethoxy)toluene (50 g, 0.24 mol) and
10 *N*-bromosuccinimide (42.75 g, 0.24 mol) in 100 mL of carbon tetrachloride under
nitrogen was added 2,2'-azobisisobutyronitrile (0.71 g, 0.004 mol). The resultant
mixture was refluxed for 2 h, then cooled to room temperature and quenched with
300 mL of water. The organic layer was collected, washed with water and brine,
dried over MgSO₄, and concentrated *in vacuo* to give 66.0 g (96%) of the desired
15 crude 3-(1,1,2,2-tetrafluoroethoxy)bromomethylbenzene product as a yellow oil.
¹H NMR indicates that this oil is a mixture of products: 7% dibrominated, 67%
monobrominated, and 20% starting material. The crude product was used
without further purification. ESMS *m/z* = 287 [M+H]⁺.

20 **EX-1B)** The crude product (56 g, 0.14 mol) from EX-1A in 200 mL of
cyclohexane was added dropwise under nitrogen to a solution of 3-phenoxyaniline
(89 g, 0.480 mol) in 500 mL of cyclohexane. The reaction mixture was refluxed
overnight, then cooled to room temperature and diluted with water and diethyl
ether. The layers were separated, and the aqueous layer was extracted with

diethyl ether. The combined organic layers were dried over MgSO_4 and concentrated *in vacuo* to give a dark oil. The crude product was purified by column chromatography on silica gel eluting with 1:4 ethyl acetate in hexane to afford 44.96 g (83%) of the desired *N*-(3-phenoxyphenyl)-[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amine product as a yellow oil. ESMS $m/z = 392$ $[\text{M}+\text{H}]^+$.

To a mixture of the amine product (15.0 g, 0.038 mol) from EX-1B and 1,1,1-trifluoro-2,3-epoxypropane (8.58 g, 0.077 mol, TCI) was added a suspension of
10 ytterbium (III) trifluoromethanesulfonate (2.37 g, 0.0031 mol) in 15 mL of acetonitrile. The resulting mixture was heated at 50 °C in a sealed glass vial for 1.5 h. The reaction mixture was cooled to room temperature then diluted with water and ethyl acetate and extracted. The organic layers were combined, dried over MgSO_4 , and concentrated *in vacuo*. The crude product was purified by
15 column chromatography on silica gel eluting with 1:4 ethyl acetate in hexane to afford 12.03 g (62%) of the desired (2*RS*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol product as a yellow oil. Anal. calcd. for $\text{C}_{24}\text{H}_{20}\text{F}_7\text{NO}_3$: C, 57.26; H, 4.00; N, 2.78. Found: C, 56.96; H, 4.35; N, 2.69. HRMS calcd. 504.1410 $[\text{M}+\text{H}]^+$, found: 504.1431.
20 ^1H NMR (CDCl_3) δ 7.28 (m, 4H), 7.14 (t, 1H), 7.07, (m, 3H), 7.00 (s, 1H), 6.94 (d, 2H), 6.46 (dd, 1H), 6.38 (dd, 1H), 6.35 (t, 1H), 5.84 (t, 1H), 4.60 (t, 2H), 4.36 (m, 1H), 3.82 (d, 1H), 3.48 (m, 1H), 2.51 (s, 1H). ^{19}F NMR (CDCl_3) δ -79.0 (s, 3F), -88.21 (d, 2F), -137.05 (dd, 2F).

EXAMPLE 2



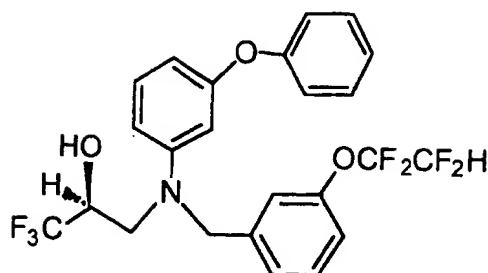
5

(2*R*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol

On a Chiralpak AD HPLC column. (2*RS*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol (12.2 g, 0.024 mol) from EX-1 was purified by chiral chromatography to give 1.4 g (0.003 mol, 12%) of (2*R*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl] amino]-1,1,1-trifluoro-2-propanol as a light yellow oil. Chiral purification was accomplished by eluting with 1:9 isopropanol in heptane at 1.0 mL/min with 250 nm UV detection. The product eluted at 8.43 min. $[\alpha]_{589} = +16.8.0$ (c 0.125 g/dL, CH₃CN), $[\alpha]_{365} = +84.0$ (c 0.125, CH₃CN). Anal. calcd. for C₂₄H₂₀F₇NO₃: C, 57.26; H, 4.00; N, 2.78. Found: C, 56.96; H, 4.35; N, 2.69. HRMS calcd.: 504.1410 [M+H]⁺, found: 504.1388. ¹H NMR (CDCl₃) δ 7.28 (m, 4H), 7.14 (t, 1H), 7.07, (m, 3H), 7.00 (s, 1H), 6.94 (d, 2H), 6.46 (dd, 1H), 6.38 (dd, 1H), 6.35 (t, 1H), 5.84 (t, 1H), 4.60 (t, 2H), 4.36 (m, 1H), 3.82 (d, 1H), 3.48 (m, 1H), 2.51 (s, 1H). ¹⁹F NMR (CDCl₃) δ -79.0 (s, 3F), -88.21 (d, 2F), -137.05 (dd, 2F).

127

EXAMPLE 3

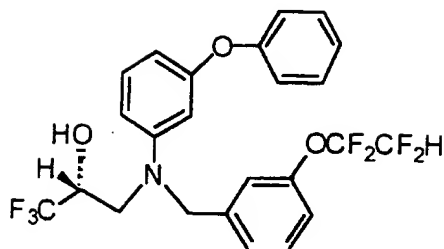


(2S)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)-
phenyl]methyl]amino]- 1,1,1-trifluoro-2-propanol

On a Chiralpak AD HPLC column. (2*RS*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol (12.2 g, 0.024 mol) from EX-1 was purified by chiral chromatography to give 10.5 g (0.021 mol, 86%) of (2*S*)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol as a light yellow oil. Chiral purification was accomplished by eluting with 1:9 isopropanol in heptane at 1.0 mL/min with 250 nm UV detection. The product eluted at 6.36 min. $[\alpha]_{589} = -17.0$ (c 0.265 g/dL, CH₃CN), $[\alpha]_{365} = -85.7$ (c 0.265, CH₃CN). Anal. calcd. For C₂₄H₂₀F₇NO₃: C, 57.26; H, 4.00; N, 2.78. Found: C, 56.96; H, 4.35; N, 2.69. HRMS calcd.: 504.1410 [M+H]⁺, found: 504.1431. ¹H NMR (CDCl₃) δ 7.28 (m, 4H), 7.14 (t, 1H), 7.07, (m, 3H), 7.00 (s, 1H), 6.94 (d, 2H), 6.46 (dd, 1H), 6.38 (dd, 1H), 6.35 (t, 1H), 5.84 (t, 1H), 4.60 (t, 2H), 4.36 (m, 1H), 3.82 (d, 1H), 3.48 (m, 1H), 2.51 (s, 1H). ¹⁹F NMR (CDCl₃) δ -79.0 (s, 3F), -88.21 (d, 2F), -137.05 (dd, 2F).

128

EXAMPLE 4



5 **(2R)-3-[(3-phenoxyphenyl)[[3-(1,1,2,2-tetrafluoroethoxy)-
phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol**

Using a procedure adopted from H.C.Brown et al. (*J. Org. Chem.* **60**, 41-46, (1995)), *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane was prepared beginning with the
10 transfer of (+)-*B*-chlorodiisopinocampheylborane ((+)-DIP-Cl, 1.2 kg, 3.74 mol) to a 5 L three neck flask containing 5 L of ether under nitrogen. Anhydrous ether (5 L) was added, and the mixture was stirred until the solids dissolved and the temperature equilibrated to 0 °C. Then 3-bromotrifluoroacetone (326 mL, 3.14 mol) was added, and the reaction was stirred for 72 h while maintaining the
15 temperature between -4 and +5 °C. The reaction was followed by ¹⁹F NMR by removing an aliquot (20 µL), quenching with anhydrous methanol (0.6 mL), and referencing to external D₂O. The reduction was 68 % complete after 48 h. The ether was removed under vacuum (100 torr to 0.1 torr), leaving a pale, viscous oil. A 5 L 3-neck flask equipped with stirrer, dropping funnel, and short-path
20 distillation head with chilled receiver was charged with 50% (w/w) aqueous NaOH and heated to 40 °C. With external heat removed, the quenched reduction mixture was added dropwise to the aqueous NaOH, with the rate controlled to maintain the pot temperature below 65 °C. The product epoxide formed immediately, distilling over with a head temperature of 32-42 °C. A yellow-

orange solid byproduct was broken up by stirring and some foaming was observed. When the distillation was complete, 145 g (43%) of the desired *R*-(+)-1,1,1-trifluoro-2,3-epoxy-propane product was obtained as a clear, colorless oil. ^1H NMR (C_6D_6) δ 2.50 (m, 1H, CF_3CH), 2.15 (dd, 1H, $J = 2.10, 5.01$ Hz), 1.75 (m, 1H). ^{19}F NMR (C_6D_6) δ -75.4 (d, $J = 4.7$ Hz). Chiral GC/MS analysis was performed on the corresponding diethylamine adduct using a gamma cyclodextrin column (Supelco gammadex120 G-cyclodextrin fused silica): 4 drops of the epoxide, *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane, and 4 drops of diethylamine were heated briefly in a sealed vial, cooled, diluted with methyl *t*-butyl ether, and analyzed. Found: two gc peaks: 10.97 min and 11.11 min (ratio 1:230; 99% ee), where the *R*-product predominated. MS calcd. for $\text{C}_7\text{H}_{14}\text{F}_3\text{NO}$: $m/z = 186$ $[\text{M}+\text{H}]^+$, found: 186, for both gc peaks. In contrast, the diethylamine adduct obtained with the TCI trifluoromethyl-oxirane (lot OGH01) from **EX-1**, gave 2 peaks with identical MS signals $m/z = 186$, 10.96 min and 11.12 min (ratio 8.5:1; 79% ee), where the *S*-product predominated.

To a mixture *N*-(3-phenoxyphenyl)-[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]-amine from **EX-1B** (1.48 g, 0.0038 mol) and *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.64 g, 0.0057 mol) was added a suspension of ytterbium (III) trifluoro-methanesulfonate (0.23 g, 0.0004 mol) in 1.5 mL of acetonitrile. The resulting mixture was heated at 50 °C in a sealed glass tube for 1.5 h. The reaction mixture was cooled to room temperature then diluted with water and ethyl acetate and extracted. The organic layers were combined, dried over MgSO_4 , and concentrated *in vacuo*. The crude product was purified by column chromatography on silica gel eluting with 1:4 ethyl acetate in hexane to afford 1.2 g (63%) of the desired (2*R*)-3-[(3-phenoxyphenyl)-[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol product as a

130

pure yellow oil (>96% ee by chiral HPLC analysis), which was identical in all respects to EX-2. Anal. calcd. for $C_{24}H_{20}F_7NO_3$: C, 57.26; H, 4.00; N, 2.78. found: C, 56.96; H, 4.35; N, 2.69. HRMS calcd.: 504.1410 $[M+H]^+$. found: 504.1431. 1H NMR ($CDCl_3$) δ 7.28 (m, 4H), 7.14 (t, 1H), 7.07, (m, 3H), 7.00 (s, 1H), 6.94 (d, 2H), 6.46 (dd, 1H), 6.38 (dd, 1H), 6.35 (t, 1H), 5.84 (t, 1H), 4.60 (t, 2H), 4.36 (m, 1H), 3.82 (d, 1H), 3.48 (m, 1H), 2.51 (s, 1H). ^{19}F NMR ($CDCl_3$) δ -79.0 (s, 3F), -88.21 (d, 2F), -137.05 (dd, 2F).

Additional examples can be prepared by one skilled in the art using similar methods and commercially available epoxides. For example, 3-[(3-phenoxyphenyl)[[3-(trifluoromethoxy)phenyl]methyl]amino]-1-chloro-2-propanols can be prepared from the reaction of *N*-(3-phenoxyphenyl)-[[3-(trifluoromethoxy)phenyl]methyl]amine with either (*R*)-epichlorohydrin or (*S*)-epichlorohydrin, as illustrated in Example Table 1.

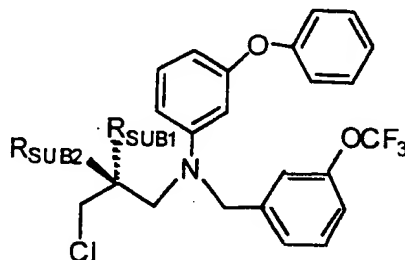
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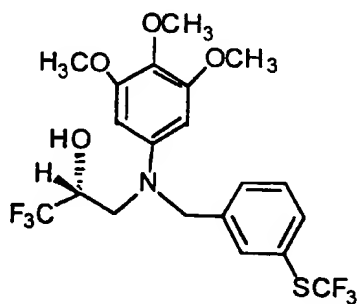
Example Table 1. 3-[(3-phenoxyphenyl)[[3-(trifluoromethoxy)phenyl]methyl]amino]-1-chloro-2-propanols.



| <u>Ex.</u> <u>No.</u> | <u>R_{SUB1}</u> | <u>R_{SUB2}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|-------------------------|-------------------------|--|--|
| 5 | OH | H | 452.1240 | 452.1245 |
| 6 | H | OH | 452.1240 | 452.1259 |

5

EXAMPLE 7



10 (2R)-3-[(3,4,5-trimethoxyphenyl)[[3-(trifluoromethylthio)phenyl]methyl]-
amino]-1,1,1-trifluoro-2-propanol

EX-45A) To a 1,2-dichloroethane (12 mL) solution of 3,4,5-trimethoxyaniline (0.80 g, 4.4 mmol) was added (3-trifluoromethylthio)benzaldehyde (0.90 g, 4.4
15 mmol), NaB(OAc)₃H (1.20 g, 5.66 mmol) and acetic acid (0.26 mL, 4.5 mmol). The cloudy solution was stirred at room temperature for 1 h. The reaction

132

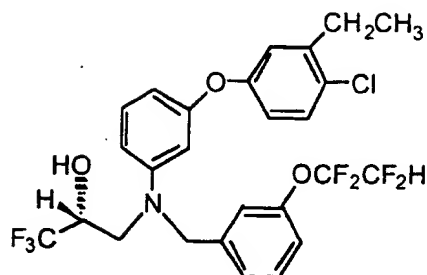
mixture was poured into water and extracted with dichloromethane. The organic layer was washed with saturated NaHCO_3 and brine, dried (MgSO_4) and evaporated to give 1.58 g (96%) of the desired *N*-(3,4,5-trimethoxyphenyl)[[3-trifluoromethylthiophenyl]methyl]amine product as an off-white solid. MS: m/z
5 $= 373.8 [\text{M}+\text{H}]^+$.

To an acetonitrile (3.2 mL) solution of amine (1.20 g, 3.2 mmol) from **EX-45A** was added *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.55 mL, 6.4 mmol) from **EX-4** and $\text{Yb}(\text{OTf})_3$ (0.40 g, 0.64 mmol). The cloudy solution was stirred in a sealed
10 flask at 50 °C for 18 h. The cooled reaction mixture was diluted with diethyl ether and washed with water and brine. The organic layer was dried (MgSO_4) and evaporated to an oil. Purification by flash column chromatography on silica gel eluting with 20% ethyl acetate in hexane gave an oil which was triturated with hexanes to give a white solid. The precipitate was isolated by filtration and dried
15 *in vacuo* to give 0.82 g (53 %) of the desired (2*R*)-3-[(3,4,5-trimethoxyphenyl)[[3-(trifluoromethylthio)phenyl] methyl]-amino]-1,1,1-trifluoro-2-propanol product as a white solid. m.p. 88.9-89.1 °C (95% ee by chiral HPLC). Anal. calcd. for $\text{C}_{20}\text{H}_{21}\text{NO}_4\text{SF}_6$: C, 49.48; H, 4.36; N, 2.89. Found: C, 49.29; H, 4.21; N, 2.81. HRMS calcd.: 486.1174 $[\text{M}+\text{H}]^+$, found: 486.1158. ^1H NMR (C_6D_6) δ 3.10 (d, 1H), 3.18 (dd, 1H), 3.32 (s, 6H), 3.53 (d, 1H), 3.64 (s, 3H), 4.01 (m, 1H), 4.21
20 (dd, 2H), 5.70 (s, 2H), 6.80 (t, 1H), 6.94 (d, 1H), 7.23 (d, 1H), 7.37 (s, 1H). $[\alpha]_{589} = +26.8$ (c 1.099 g/dL, CHCl_3).

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133

EXAMPLE 8



5 **(2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol**

EX-8A) To a solution of 1,3-dinitrobenzene (16.8 g, 0.1 mol) and 4-chloro-3-ethylphenol (15.6 g, 0.1 mol) in 200 mL of dimethylsulfoxide was added cesium carbonate (65 g, 0.2 mol). The reaction mixture was heated at 100 °C under nitrogen overnight then cooled to room temperature. The reaction mixture was filtered through celite then rinsed with diethyl ether and a small amount of water. The filtrate was extracted several times with diethyl ether. The organic layers were combined, washed with water and brine, dried over MgSO₄, and concentrated *in vacuo* to give 21.8 g (78%) of the desired 3-(4-chloro-3-ethylphenoxy)-1-nitrobenzene product as a dark orange oil, which was greater than 90% pure by reverse phase HPLC analysis. HRMS calcd. for C₁₄H₁₂ClNO₃: 295.0849 [M+NH₄]⁺, found 295.0862.

EX-8B) To a solution of 3-(4-chloro-3-ethylphenoxy)-1-nitrobenzene (10 g, 0.036 mol) from EX-8A in 400 mL of glacial acetic acid and 1 mL of water was added zinc metal (20 g, 0.305 mol) at room temperature, and the resultant mixture was stirred for 1 h. The reaction mixture was filtered through celite. The filtrate was neutralized with ammonium hydroxide and extracted with diethyl ether. The

organic layer was washed with water and brine, dried over MgSO_4 , and concentrated *in vacuo* to give 10 g (100%) of the desired 3-(4-chloro-3-ethylphenoxy)aniline product as a dark orange oil, which was greater than 90% pure by reverse phase HPLC analysis. HRMS calcd. for $\text{C}_{14}\text{H}_{14}\text{ClNO}$: 248.0842

5 $[\text{M}+\text{H}]^+$, found: 248.0833.

EX-8C) To a solution of 3-(4-chloro-3-ethylphenoxy)aniline (2.0 g, 8.1 mmol) from **EX-8B** and 3-(1,1,2,2-tetrafluoroethoxy)benzaldehyde (1.6 g, 7.3 mmol) in 30 mL of dichloroethane was added sodium triacetoxymethylborohydride (2.0 g, 9.7

10 mmol) and glacial acetic acid (0.51 mL, 8.9 mmol). The reaction mixture was stirred at room temperature for 1 h then quenched with water and extracted with diethyl ether. The organic layer was washed with water and brine, dried over MgSO_4 , and concentrated *in vacuo* to give 3.5 g (95%) of the desired *N*-[(4-chloro-3-ethylphenoxy)phenyl]-3-[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]

15 amine product as a brown oil, which was greater than 90% pure by reverse phase HPLC analysis. HRMS calcd. for $\text{C}_{23}\text{H}_{20}\text{ClF}_4\text{NO}_2$: 454.1197 $[\text{M}+\text{H}]^+$, found: 454.1220.

A solution of *N*-[(4-chloro-3-ethylphenoxy)phenyl]-3-[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amine (1.8 g, 4.0 mmol) from **EX-8C**, *R*-(+)-1,1,1-trifluoro-2,3-epoxy-propane (0.64 g, 0.0057 mol) from **EX-4**, and ytterbium (III) trifluoromethanesulfonate (0.25 g, 0.4 mmol) in 1.5 mL of acetonitrile was

20 heated at 40 °C in a sealed glass tube for 1 h. The reaction mixture was cooled to room temperature then diluted with water and diethyl ether and extracted. The

25 ether layer was washed with water and brine, dried over MgSO_4 , and concentrated *in vacuo*. The crude product was purified by column

135

chromatography on silica gel eluting with 1:7:0.01 of ethyl acetate:hexane:ammonium hydroxide to afford 1.5 g (66%) of the desired (2*R*)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl]][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-tri-fluoro-2-propanol product as a yellow oil (96% ee by
5 chiral HPLC analysis). $[\alpha]_{589}^{25} = +36.9$ (c 1.044g%, CHCl₃). $[\alpha]_{365}^{25} = +189.7$ (c 1.044g%, CHCl₃). The refractive index @ 25 °C is 1.5275. Anal. calcd. for C₂₆H₂₃ClF₇NO₃: C, 55.18; H, 4.10; N, 2.48. found: C, 54.92; H, 4.05; N, 2.33. HRMS calcd.: 566.1330 [M+H]⁺, found: 566.1323. ¹H NMR (CDCl₃) δ 7.30 (t, 1H), 7.20 (d, 1H), 7.15 (t, 1H), 7.08 (t, 2H), 7.00 (s, 1H), 6.86 (d, 1H),
10 6.68 (dd, 1H), 6.48 (dd, 1H), 6.36 (dd, 1H), 6.34 (t, 1H), 5.81 (tt, 1H), 4.62 (s, 2H), 4.32 (m, 1H), 3.84 (dd, 1H), 3.55 (dd, 1H), 2.67 (q, 2H), 2.45 (bs, 1H), 1.17 (t, 3H). ¹⁹F NMR (CDCl₃) δ -79.22 (d, 3F), -88.57 (m, 2F), -137.16 (dt, 2F).

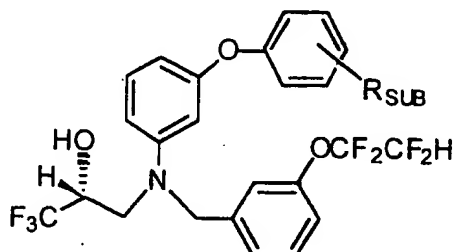
Additional examples of (2*R*)-3-[[3-(substituted-phenoxy)phenyl]][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols and
15 (2*R*)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl]][[3-substituted-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols can be prepared by one skilled in the art using similar methods, as shown in Example Tables 2 and 3, respectively.

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136

Example Table 2. (2*R*)-3-[[3-(Substituted-phenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols.



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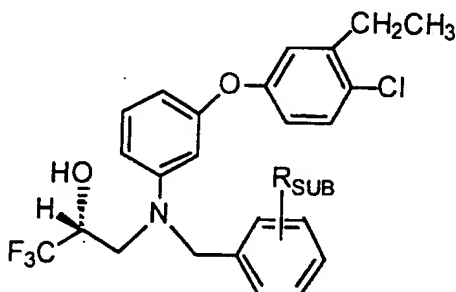
| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|------------------------|--|--|
| 9 | 4-methyl | 518.1566 | 518.1587 |
| 10 | 3-isopropyl | 546.1879 | 546.1900 |
| 11 | 3-ethyl | 532.1723 | 532.1713 |

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Example Table 3. (2*R*)-3-[[3-(4-Chloro-3-ethylphenoxy)phenyl]][[3-substituted-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols.



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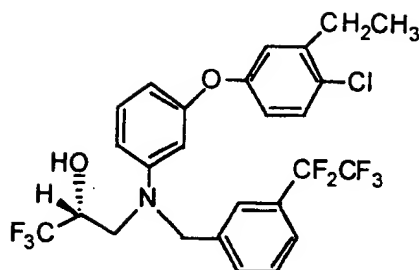
| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|-----------------------------|--|--|
| 12 | 3-trifluoromethoxy | 534.1271 | 534.1309 |
| 13 | 3-trifluoromethyl, 4-fluoro | 536.1228 | 536.1265 |
| 14 | 2-fluoro, 4-trifluoromethyl | 536.1228 | 536.1241 |
| 15 | 2-trifluoromethyl, 4-fluoro | 536.1228 | 536.1245 |
| 16 | 2-fluoro, 5-trifluoromethyl | 536.1228 | 536.1252 |
| 17 | 2-fluoro, 6-trifluoromethyl | 536.1228 | 536.1199 |

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138

EXAMPLE 18



(2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(1,1,1,2,2-pentafluoroethyl)-
 5 phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol

EX-18A) Sodium pentafluoroethyl propionate (8.4 g, 50 mmol) and 3-iodotoluene (5.5 g, 25 mmol) were dissolved in anhydrous DMF (300 mL) under nitrogen. CuI (9.5 g, 50 mmol) was added, and the mixture was heated to 160 °C under
 10 nitrogen for 4 h, at which time a 15 mL fraction of a mixture of DMF and 3-pentafluoroethyl toluene was collected. The distillate was diluted with Et₂O and was washed with brine. The ether layer was dried over MgSO₄, filtered and concentrated *in vacuo* to give 5.25 g (55%) of the desired 3-pentafluoroethyl-toluene product as a colorless oil. ¹H NMR (CDCl₃) δ 7.36 (m, 4H), 2.40 (s,
 15 3H). ¹⁹F NMR (CDCl₃) δ -85.2 (s, 3F), -115.2 (s, 2F).

EX-18B) The 3-pentafluoroethyl-toluene (2.9 g, 13.8 mmol) product from EX-18A and N-bromosuccinimide (2.5 g, 13.8 mmol) were dissolved in CCl₄ (25 mL). AIBN (50 mg, 0.3 mmol) was added, and the mixture was refluxed for 3.5 h under
 20 N₂. The reaction mixture was cooled to room temperature and diluted with water. The layers were separated, and the organic layer was washed with brine, dried with anhydrous MgSO₄, filtered, and concentrated *in vacuo* to give 3.4 g

(87%) of a colorless oil. The ^1H NMR spectrum indicated that the crude product contained 3-pentafluoroethyl-benzylbromide (70%), the benzyldibromide (10%) and pentafluoroethyl toluene (20%). ^1H NMR (CDCl_3) δ 7.60 (m, 2H), 7.50 (m, 2H), 4.50 (s, 2H). ^{19}F NMR (CDCl_3) δ -85.1 (s, 3F), -115.4 (s, 2F).

5

EX-18C) A solution of 3-(4-chloro-3-ethylphenoxy)aniline (1.7 g, 6.9 mmol) was prepared in cyclohexane (13 mL). A solution of crude 3-pentafluoroethyl benzylbromide (1 g, 3.5 mmol) product from **EX-18B** in cyclohexane (10 mL) was added dropwise under nitrogen over 3 min. The reaction mixture was refluxed under N_2 for 24 h and then was cooled to room temperature. The mixture was diluted with Et_2O and saturated aqueous NaHCO_3 . The layers were separated, and the aqueous layer was extracted with Et_2O . The organic layer was washed with brine, dried with anhydrous MgSO_4 , filtered and concentrated *in vacuo*. The residue was purified by chromatography on silica gel eluting with hexanes in ethyl acetate (95:5) which gave 0.56 g (35%) of the desired *N*-[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(pentafluoro-ethyl)phenyl]methyl]amine product as a brown oil. ^1H NMR (CDCl_3) δ 7.53 (m, 4H), 7.27 (d, 1H), 7.15 (t, 1H), 6.93 (d, 1H), 6.77 (dd, 1H), 6.41 (tt, 2H), 6.30 (t, 1H), 4.41 (s, 2H), 2.73 (q, 2H), 1.23 (t, 3H). ^{13}C NMR (CDCl_3) δ 158.6, 156.1, 143.4, 141.3, 140.2, 131.3, 130.7, 130.4, 129.4, 128.1, 120.4, 117.8, 108.8, 103.9, 48.5, 27.5, 14.1. ^{19}F NMR (CDCl_3) δ -85.1 (s, 3F), -115.2 (s, 2F). HRMS calcd. for $\text{C}_{23}\text{H}_{19}\text{ClF}_5\text{NO}$: 456.1154 $[\text{M}+\text{H}]^+$, found: 456.1164.

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The *N*-[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amine (0.4 g, 0.88 mmol) product of **EX-18C** was dissolved in anhydrous

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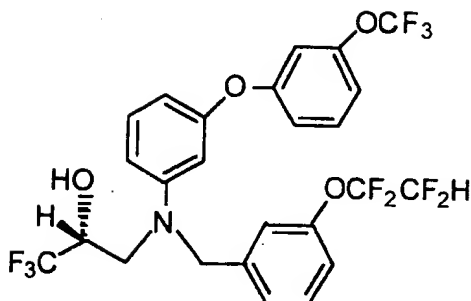
acetonitrile (1.5 mL). *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.22 g, 1.94 mmol) and Yb(OTf)₃ (22 mg, 0.035 mmol) were added, and the reaction mixture was stirred under N₂ at 45 °C in a sealed glass tube for 15 h. The reaction mixture was then cooled to room temperature and diluted with Et₂O and saturated aqueous NaHCO₃. The layers were separated and the aqueous layer was extracted with Et₂O. The ether layers were combined, washed with brine, dried with anhydrous Na₂SO₄, filtered, and concentrated *in vacuo*. The viscous oil was adsorbed onto silica gel and purified by column chromatography eluting with hexanes in ethyl acetate (95:5) which gave 0.32 g (64%) of the desired (2*R*)-3-[(4-chloro-3-ethylphenoxy)phenyl][3-(pentafluoroethyl)phenyl]-methylamino]-1,1,1-trifluoro-2-propanol product as a viscous, colorless oil. ¹H NMR (CDCl₃) δ 7.47 (m, 4H), 7.23 (m, 3H), 6.90 (d, 1H), 6.72 (dd, 1H), 6.52 (d, 1H), 6.42 (m, 2H), 4.73 (s, 2H), 4.39 (m, 1H), 3.91 (dd, 1H), 3.58 (m, 2H), 2.73 (q, 2H), 2.57 (s, 1H), 1.22 (t, 3H). ¹⁹F NMR (CDCl₃) δ -79.2 (s, 3F), -84.9 (s, 3F), -115.2 (s, 2F). HRMS calcd. for C₂₆H₂₂ClF₈NO₂: 568.1290 [M+H]⁺, found: 568.1296.

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EXAMPLE 19



5 **(2R)-3-[[3-(3-trifluoromethoxyphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol**

EX-19A) To a solution of 1,3-dinitrobenzene (4.5 g, 0.03 mol) and 3-trifluoromethoxy-phenol (4.8 g, 0.03 mol) in 54 mL of dimethylsulfoxide was added cesium carbonate (21.8 g, 0.07 mol). The reaction mixture was heated at
 10 100 °C under nitrogen overnight then cooled to room temperature. The reaction mixture was diluted with water and extracted with diethyl ether several times. The organic layers were combined, washed with 1 N HCl and water, dried over MgSO₄, and concentrated *in vacuo*. The crude product was purified by column chromatography on silica gel eluting with 1:9 ethyl acetate in hexane to afford 3.0
 15 g (38%) of the desired 3-(3-trifluoro-methoxyphenoxy)nitrobenzene product as a yellow-orange liquid which was 85% pure by reverse phase HPLC analysis. This material was carried on without further purification.

EX-19B) To a solution of 3-(3-trifluoromethoxyphenoxy)nitrobenzene (3.0 g, 0.01 mol) from EX-19A in 100 mL of glacial acetic acid was added zinc metal (6.6 g, 0.1 mol) at room temperature, and the resultant mixture was stirred for 1 h. The reaction mixture was filtered through celite. The filtrate was neutralized with ammonium hydroxide and extracted with diethyl ether then ethyl acetate. The
 20

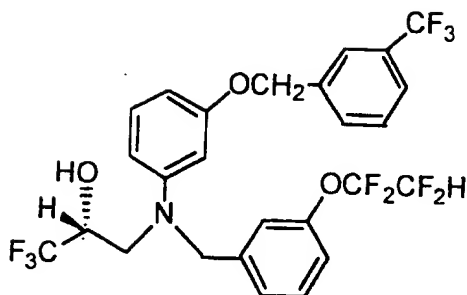
combined organic layers were dried over MgSO_4 and concentrated *in vacuo*. The crude product was purified by column chromatography on silica gel eluting with 1:9 ethyl acetate in hexane to afford 1.2 g (44%) of the desired 3-(3-trifluoromethoxyphenoxy)aniline product as a yellow oil which was 98% pure by reverse phase HPLC analysis. Anal. calcd. for $\text{C}_{13}\text{H}_{10}\text{F}_3\text{NO}_2$: C, 58.00; H, 3.74; N, 5.20. found: C, 57.68; H, 3.57; N, 5.14. HRMS calcd.: 270.0742 $[\text{M}+\text{H}]^+$, found: 270.0767.

EX-19C) To a solution of 3-(3-trifluoromethoxyphenoxy)aniline (1.0 g, 3.7 mmol) from **EX-19B** and 3-(1,1,2,2-tetrafluoroethoxy)benzaldehyde (0.83 g, 3.7 mmol) in 18.5 mL of dichloroethane was added sodium triacetoxyborohydride (1.0 g, 4.7 mmol) and glacial acetic acid (0.25 mL, 4.3 mmol). The reaction mixture was stirred at room temperature overnight then quenched with saturated aqueous sodium bicarbonate and extracted with methylene chloride. The organic layer was dried over MgSO_4 and concentrated *in vacuo* to give 1.8 g (100%) of the desired [3-(3-trifluoromethoxy-phenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl] amine product as a yellow oil. which was greater than 90% pure by reverse phase HPLC analysis. HRMS calcd. for $\text{C}_{22}\text{H}_{16}\text{F}_7\text{NO}_3$: 476.1097 $[\text{M}+\text{H}]^+$, found: 476.1069. This material was carried on to the next step without further purification.

A solution of [3-(3-trifluoromethoxyphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]methyl]amine (1.8 g, 3.7 mmol) from **EX-19C**, *R*-(+)-1,1,1-trifluoro-2,3-epoxy-propane (0.57 g, 5.2 mmol), and ytterbium (III) trifluoromethanesulfonate (0.24 g, 0.38 mmol) in 2.0 mL of acetonitrile was heated at 40 °C in a sealed glass tube overnight. At this time reverse phase HPLC analysis indicated that the reaction was only 50% complete. Additional

- ytterbium (III) trifluoromethanesulfonate and *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.26 g, 2.3 mmol) were added to the reaction mixture and again heated at 40 °C in a sealed glass tube for 48 h. The reaction mixture was cooled to room temperature then diluted with water and methylene chloride and extracted.
- 5 The organic layer was washed with brine, dried over MgSO₄, and concentrated *in vacuo*. The crude product was purified by reverse phase HPLC eluting with 30% to 90% acetonitrile in water to afford 1.25 g (23%) of the desired (2*R*)-3-[[3-(3-trifluoromethoxyphenoxy) phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol product as yellow-brown oil (90% ee
- 10 by chiral HPLC analysis). HRMS calcd. for C₂₅H₁₉F₁₀NO₄: 588.1233 [M+H]⁺, found: 588.1225. ¹H NMR (CDCl₃) δ 7.35-7.18 (m, 3H), 7.12 (t, 2H), 7.01 (s, 1H), 6.93 (d, 1H), 6.85 (d, 1H), 6.82 (s, 1H), 6.56 (dd, 1H), 6.47 (dd, 1H), 6.41 (s, 1H), 5.88 (t, 1H), 4.66 (s, 2H), 4.35 (m, 1H), 3.86 (d, 1H), 3.59 (dd, 1H), 2.02 (s, 1H). ¹⁹F NMR (CDCl₃) δ -58.31 (s, 3F), -79.24 (d, 3F), -
- 15 88.57 (m, 2F), -137.16 (dt, 2F).

EXAMPLE 20



5 (2R)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl][3-[(trifluoromethyl)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol

EX-20A) To a solution of 3-aminophenol (4.91 g, 45.0 mmol) and 3-(1,1,2,2-tetrafluoroethoxy)benzaldehyde (10.0 g, 45.0 mmol) in 100 mL of 1,2-dichloroethane was added sodium triacetoxyborohydride (14.28 g 67.5 mmol) and glacial acetic acid (2.7 mL, 47.3 mmol). The reaction mixture was stirred at room temperature for 6 h then quenched with water and extracted with dichloromethane. The organic layer was washed with saturated aqueous sodium bicarbonate, dried over MgSO₄, and concentrated *in vacuo* to give 11.82 g (83%) of the desired 3-[[[3-(1,1,2,2-tetrafluoro-ethoxy)phenyl]methyl]amino]phenol product as a dark orange oil. ¹H NMR (acetone-*d*₆) δ 7.01-7.38 (m, 5H), 6.26-6.44 (m, 3H), 6.08 (t, 1H), 5.88 (tt, 1H), 4.34 (s, 2H).

20 EX-20B) A solution of 3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]phenol (5.1 g, 16.2 mmol) from EX-20A, *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (1.5 mL, 17.4 mmol), and ytterbium trifluoromethanesulfonate (1.0 g, 10 mol%) in 10 mL of acetonitrile was heated at 50 °C in a sealed glass tube for 4 h. The

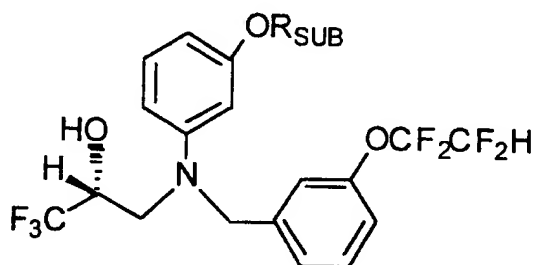
reaction mixture was cooled to room temperature, then diluted with water and diethyl ether and extracted. The ether layer was washed with saturated aqueous sodium bicarbonate and brine, dried over MgSO_4 , and concentrated *in vacuo* to give 5.64 g (81%) of the desired (2*R*)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl][3,3,3-trifluoro-2-hydroxy-propyl)amino]phenol product as a yellow oil. ^1H NMR (acetone- d_6) δ 7.41 (t, 1H), 7.23 (d, 1H), 7.16-7.20 (m, 2H), 6.97 (t, 1H), 6.42 (tt, 1H), 6.18-6.24 (m, 3H), 4.77 (s, 2H), 4.43-4.48 (m, 1H), 3.58 (dd, 1H), 3.39 (dd, 1H).

- 10 To a solution of (2*R*)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl][3,3,3-trifluoro-2-hydroxypropyl)amino]phenol (100 mg, 0.23 mmol) from EX-20B and 3-trifluoromethylbenzyl bromide (70.0 mg, 0.27 mmol) in 2.5 mL of acetone was added cesium carbonate (100 mg, 0.31 mmol). The reaction mixture was heated at 60 °C for 18 h then cooled to room temperature. The reaction mixture was
- 15 filtered through celite, and the filtrate was concentrated. The residue was purified by reverse phase HPLC eluting with 50% to 90% acetonitrile in water to afford 63.3 mg (45%) of the desired (2*R*)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl][3-[[3-(trifluoro-methyl)phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol product as an orange oil.
- 20 HRMS calcd. for $\text{C}_{26}\text{H}_{21}\text{F}_{10}\text{NO}_3$: 586.1440 $[\text{M}+\text{H}]^+$, found: 586.1419. ^1H NMR (acetone- d_6) δ 7.61-7.82 (m, 4H), 7.41 (t, 1H), 7.25 (d, 1H), 7.10-7.21 (m, 3H), 6.34-6.67 (m, 4H), 5.73 (d, 1H), 5.19 (s, 2H), 4.82 (s, 2H), 4.34-4.48 (m, 1H), 3.99 (dd, 1H), 3.68 (dd, 1H).

Additional examples of (2*R*)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]-[3-[[aryl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanols are prepared by one skilled in the art using similar methods, as shown in. Example Table 4.

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Example Table 4. (2*R*)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl][3-[[aryl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanols.

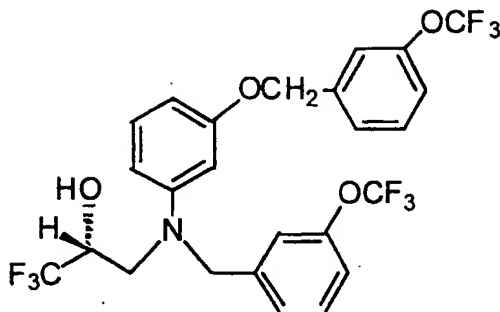


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| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|--------------------------|--|--|
| 21 | 3,5-difluorobenzyl | 554.1378 | 554.1352 |
| 22 | 3-trifluoromethoxybenzyl | 602.1389 | 602.1390 |
| 23 | 3-isopropyl | 470.1566 | 464.1601 |

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EXAMPLE 24



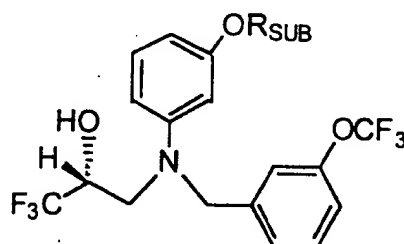
5 **(2*R*)-3-[[[3-[[3-(trifluoromethoxy)phenyl]methoxy]phenyl][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol**

(2*R*)-3-[[[3-[[3-(trifluoromethoxy)phenyl]methoxy]phenyl][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol can be prepared by one skilled in the art using similar methods starting from 3-(trifluoromethoxy)-
 10 benzaldehyde. HRMS calcd. for C₂₅H₂₀F₉NO₄: 570.1327 [M+H]⁺, found: 570.1325. ¹H NMR (acetone-*d*₆) δ 7.43 (t, 1H), 7.32 (d, 1H), 7.18-7.23 (m, 2H), 7.01-7.16 (m, 3H), 6.92-7.00 (m, 1H), 6.38-6.45 (m, 3H), 5.12 (s, 2H), 4.81 (s, 2H), 4.41-4.53 (m, 1H), 3.98 (dd, 1H), 3.63 (dd, 1H).

15 Additional examples of (2*R*)-3-[[[3-[[aryl]methoxy]phenyl][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols can be prepared by one skilled in the art using similar methods, as shown in Example Table 5.

148

Example Table 5. (2*R*)- 3-[[3-[[aryl]methoxy]phenyl][[3-(trifluoromethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols.



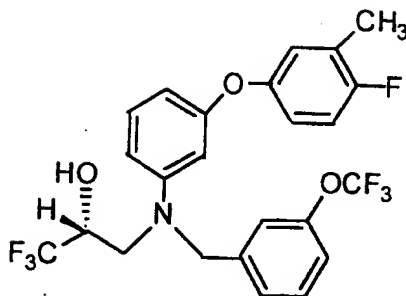
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| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|-----------------------------------|--|--|
| 25 | 4-trifluoromethoxybenzyl | 570.1327 | 570.1299 |
| 26 | 3,5- di(trifluoromethyl)benzyl | 622.1252 | 622.1252 |
| 27 | 3-trifluoromethylbenzyl | 554.1378 | 554.1369 |
| 28 | 3,5-difluorobenzyl | 522.1315 | 522.1259 |
| 29 | benzyl | 486.1504 | 486.1504 |
| 30 | isopropyl | 438.1504 | 438.1509 |
| 31 | cyclohexylmethyl | 492.1973 | 492.1973 |
| 32 | cyclopentyl | 464.1660 | 464.1641 |

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EXAMPLE 33



5 **(2R)-3-[[3-(4-fluoro-3-methylphenoxy)phenyl][[3-(trifluoromethoxy)-
phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol**

EX-33A) To a solution of 3-bromoaniline (5.7 mL, 52.6 mmol) and 3-trifluoro-
methoxybenzaldehyde (10.0 g, 52.6 mmol) in 135 mL of dichloroethane was
10 added sodium triacetoxyborohydride (14.5 g, 68.4 mmol) and glacial acetic acid
(3.1 mL, 54.7 mmol). The reaction was stirred at room temperature for 2 h, then
quenched with water and extracted with dichloromethane. The organic layer was
washed with saturated aqueous sodium bicarbonate, dried over MgSO₄, and
concentrated *in vacuo*. The crude product was purified by column
15 chromatography on silica gel eluting with 1:9 ethyl acetate in hexane to give 14.3 g
(78%) of the desired of *N*-(3-bromophenyl)[[3-(trifluoromethoxy)
phenyl]methyl]amine product as a dark brown oil. HRMS calcd. for
C₁₄H₁₁BrF₃NO: 346.0055 [M+H]⁺, found: 346.0052.

20 **EX-33B)** A solution of of *N*-(3-bromophenyl)[[3-
(trifluoromethoxy)phenyl]methyl]-amine (10.0 g, 28.9 mmol) from EX-33A, *R*-
(+)-1,1,1-trifluoro-2,3-epoxypropane (4.2 g, 37.6 mmol), and ytterbium (III)
trifluoromethanesulfonate (1.79 g, 2.89 mmol) in 27 mL of acetonitrile was heated

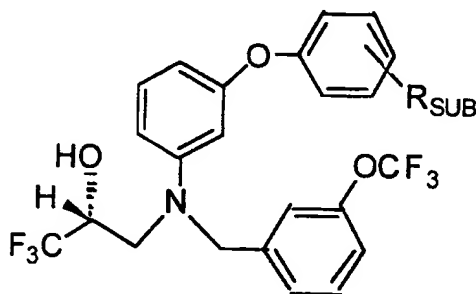
at 50 °C in a sealed glass tube overnight. The reaction mixture was cooled to room temperature and filtered through celite. The crude product was purified by column chromatography on silica gel eluting with 2:3 dichloromethane in hexane to afford 11.9 g (90%) of the desired (2*R*)-3-[[[3-(bromophenyl)][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol product as a brown oil (98% ee by chiral HPLC analysis). HRMS calcd. for $C_{17}H_{14}BrF_6NO_2$: 458.0190 $[M+H]^+$, found: 458.0197.

A suspension of 4-fluoro-3-methylphenol (98.0 μ L, 0.88 mmol) and cesium carbonate (319.5 mg, 0.98 mmol) in 1 mL of *N,N*-dimethylacetamide was preheated at 60 °C for 5 minutes. To this solution was added 4 mL of a stock solution containing (2*R*)-3-[[[3-(bromophenyl)][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol (200 mg, 0.437 mmol) from EX-33B, 1-naphthoic acid (164 mg, 0.95 mmol), copper(I) trifluoromethanesulfonate benzene complex (21.8 mg, 0.0434 mmol), 4 Å sieves (105 mg), and 4 mL of toluene. The reaction mixture was stirred at 105 °C for 3 weeks and 2 days. During that time, additional cesium carbonate and catalyst were added (a spatula tip of each) to the reaction three different times. The reaction was cooled to room temperature, filtered through celite, and the solvent was evaporated. The residue was purified by reverse phase HPLC eluting with 35% to 90% acetonitrile in water to afford 50.5 mg (23%) of the desired (2*R*)-3-[[[3-(4-fluoro-3-methylphenoxy)phenyl][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol product as an orange oil. HRMS calcd. for $C_{24}H_{20}F_7NO_3$: 504.1410 $[M+H]^+$, found: 504.1389. 1H NMR (acetone- d_6) δ 7.44 (t, 1H), 7.24 (d, 1H), 7.08-7.21 (m, 3H), 6.98 (t, 1H), 6.75-6.85 (m, 1H), 6.68-6.74 (m, 1H), 6.53 (d, 1H), 6.21-6.34 (m, 2H), 4.79 (t, 2H), 4.46-4.53 (m, 1H), 3.95 (dd, 1H), 2.61-2.72 (m, 1H), 2.20 (s, 3H).

Additional examples (2*R*)-3-[[*(aryloxy)phenyl*][[3-(trifluoromethoxy)*phenyl*] methyl]amino]-1,1,1-trifluoro-2-propanols can be prepared by one skilled in the art using similar methods, as shown in Example Table 6.

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Example Table 6. (2*R*)-3-[[*(aryloxy)phenyl*][[3-(trifluoromethoxy)*phenyl*]-methyl]amino]-1,1,1-trifluoro-2-propanols.



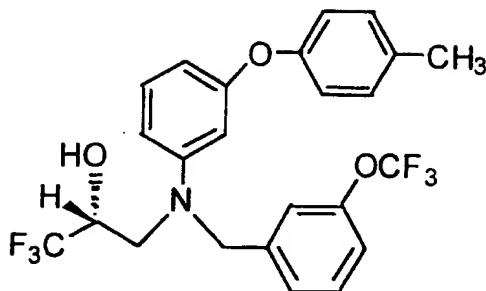
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| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|---|--|--|
| 34 | 3-trifluoromethoxy | 556.1170 | 556.1180 |
| 35 | 3-isopropyl | 514.1817 | 514.1823 |
| 36 | 3,4-dimethyl | 500.1660 | 500.1654 |
| 37 | 4-chloro-3-methyl | 520.1114 | 520.1129 |
| 38 | 3- <i>tert</i> -butyl | 528.1973 | 528.1942 |
| 39 | 3,4-dichloro | 540.0568 | 540.0567 |
| 40 | 3,4-(CH ₂ CH ₂ CH ₂ CH ₂)- | 526.1817 | 526.1788 |

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152

EXAMPLE 41



(2R)-3-[[[3-(4-methylphenoxy)phenyl]][[3-(trifluoromethoxy)phenyl]
methyl]amino]- 1,1,1-trifluoro-2-propanol

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EX-41A) To a solution of *p*-cresol (5.76 g, 0.053 mol) and 1,3-dinitrobenzene (8.97 g, 0.053 mol) in 100 mL of dimethylsulfoxide was added cesium carbonate (43.4 g, 0.133 mol). The reaction mixture was heated at 100 °C for 18 h, then
10 cooled to room temperature, quenched with water, and extracted with diethyl ether. The organic layers were combined, washed with 0.1 N HCl and water, dried over MgSO₄ and concentrated *in vacuo*. The crude product was purified by column chromatography on silica gel eluting with 1:4 ethyl acetate in hexane to afford 8.0 g (66%) of the desired 3-(4-methylphenoxy)nitrobenzene product as a
15 yellow oil. ¹H NMR (CDCl₃) δ 7.83 (s, 1H), 7.64 (t, 1H), 7.32 (d, 1H), 7.18 (d, 1H), 7.09 (d, 2H), 6.8 (d, 2H), 2.20 (s, 1H).

EX-41B) A solution of 3-(4-methylphenoxy)nitrobenzene (8.0 g, 0.035 mol) from EX-41A in 25 mL of ethanol under nitrogen was charged with 10% palladium on carbon (0.80 g). The resulting mixture was hydrogenated for 4 h at room
20 temperature and 45 psi. The reaction mixture was filtered through celite and concentrated *in vacuo* to give 6.7 g (96%) of the desired 3-(4-methylphenoxy)aniline product as a yellow oil. ESMS *m/z* = 200

$[M+H]^+$ confirmed the desired $C_{13}H_{13}NO$ product and the complete consumption of starting material.

- EX-41C)** To a solution of 3-(4-methylphenoxy)aniline (2.91 g, 0.015 mol) from **EX-41B**, and 3-(trifluoromethoxy)benzaldehyde (3.24 g, 0.015 mol) in 50 mL dichloroethane was added sodium triacetoxyborohydride (4.02 g, 0.019 mol) and glacial acetic acid (0.99 g, 0.017 mol). The reaction mixture was stirred at room temperature for 18 h, then quenched with saturated aqueous sodium bicarbonate and extracted with dichloromethane. The organic layers were combined, dried over $MgSO_4$ and concentrated *in vacuo* to give 5.38 g (91%) of the desired *N*-[3-(4-methylphenoxy)-phenyl]-[[3-(trifluoromethoxy)phenyl]methyl]amine product as an orange oil. ESMS $m/z = 374$ $[M+H]^+$ confirmed the desired $C_{21}H_{18}NO_2F_3$ product and the complete consumption of starting material.
- To a mixture of *N*-[3-(4-methylphenoxy)phenyl]-[[3-(trifluoromethoxy)phenyl]methyl]amine (1.3 g, 0.0035 mol) from **EX-41C** and *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.59 g, 0.0053 mol) was added a suspension of ytterbium (III) trifluoromethanesulfonate (0.22 g, 0.0004 mol) in 1.3 mL of acetonitrile. The resulting mixture was heated at 50 °C in a sealed glass tube for 18 h. The reaction mixture was cooled to room temperature, then diluted with water and extracted with ethyl acetate. The crude product was purified by column chromatography on silica gel eluting with 1:4 ethyl acetate in hexane to afford 1.03 g (61%) of the desired (2*R*)-3-[3-(4-methyl-phenoxy)phenyl][[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propa-nol product as a pure yellow oil. Anal. calcd. for $C_{24}H_{21}F_6NO_3$: C, 59.38; H, 4.36; N, 2.89. Found: C, 59.17; H, 4.62; N, 2.80. HRMS calcd.: 486.1504 $[M+H]^+$, found: 486.1513. 1H NMR (C_6D_6) δ 6.82 (m, 8H), 6.60 (dd, 1H), 6.42 (dd, 1H), 6.38 (s, 1H), 6.18 (dd, 1H), 4.00 (s,

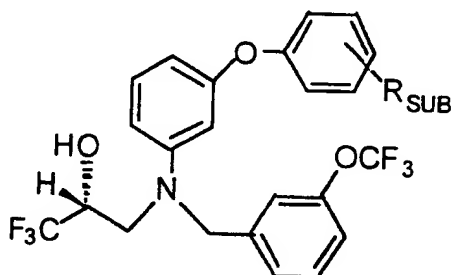
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2H), 3.63 (m, 1H), 3.40 (d, 1H), 3.02 (m, 1H), 2.00 (s, 3H), 1.40 (d, 1H). ^{19}F

NMR (C_6D_6) δ -57.98 (s, 3F), -78.50 (s, 3F).

Additional examples of (2*R*)-3-[3-(substituted-phenoxy)phenyl]-[[3-(trifluoro-methoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols can
 5 prepared by one skilled in the art using similar methods, as shown in Example Table 7.

Example Table 7. (2*R*)- 3-[3-(substituted-phenoxy)phenyl][[3-(trifluoromethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanols.
 10

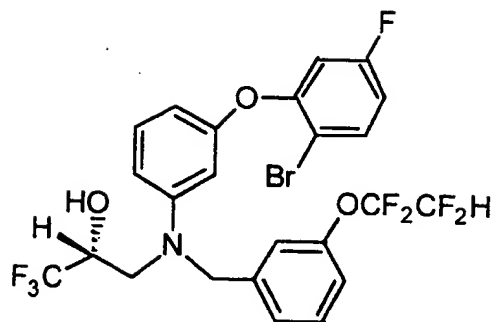


| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|------------------------|--|--|
| 42 | 4-fluoro | 490.1253 | 490.1238 |

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EXAMPLE 43



5

(2*R*)-3-[[[3-(2-bromo-5-fluorophenoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol

EX-43A) To a solution of 3-aminophenol (5 g, 46 mmol), 1-bromo-2,4-difluoro-
 10 benzene (10 g, 50 mmol) and Cs_2CO_3 (16 g, 50 mmol) in 25 mL of dimethyl-
 formamide was added solid $(\text{CuOTf})_2\text{C}_6\text{H}_6$ (100 mg), and the mixture was stirred
 under nitrogen at 85 °C for 22 h, at which time HPLC analysis indicated that the
 reaction had gone to completion and formed two products. The DMF was
 removed under reduced pressure. The residue was diluted with ether and filtered
 15 through a celite pad. The pad was washed with ether and a small amount of water.
 The mixture was extracted with ether several times. The combined ether layers
 were washed with water and brine, then dried over MgSO_4 . The dried organic
 layer was evaporated to give 10.2 g (80%) of the desired product, which consisted
 of a 11:1 ratio of 3-(2-bromo-5-fluoro-phenoxy)aniline and 3-(4-bromo-3-
 20 fluorophenoxy)aniline. The crude product was purified by flash column
 chromatography on silica gel eluting with 1:7:0.01 of ethyl
 acetate:hexane:ammonium hydroxide to give 8.8 g (68%) of the desired product as

a yellow oil, which was a 25:1 ratio of 3-(2-bromo-5-fluorophenoxy)aniline and 3-(4-bromo-3-fluorophenoxy)aniline. HRMS calcd. for $C_{12}H_9NOFBr$: 281.9930 $[M+H]^+$, found: 281.9950.

- 5 **EX-43B)** The 3-(2-bromo-5-fluorophenoxy)aniline (1.39 g, 4.95 mmol) product from **EX-43A** and 3-(1,1,2,2-tetrafluoroethoxy)benzaldehyde (1.0 g, 4.5 mmol) were dissolved in 15 mL of dichloroethane and acetic acid (0.30 mL, 5.4 mmol), then solid $NaBH(OAc)_3$ (1.26 g, 5.9 mmol) was added. The mixture was stirred at room temperature for 1 h, then quenched with water and extracted with ether.
- 10 The ether layer was washed with water and brine, then dried over $MgSO_4$, and evaporated to give 2.1 g (97%) of crude product, which was purified by flash column chromatography on silica gel eluting with 1:7:0.01 of ethyl acetate:hexane:ammonium hydroxide to give 2.0 g (91%) of the desired 3-[3-(2-bromo-5-fluoro-phenoxy)phenyl][[3-(1,1,2,2-tetrafluoro-
- 15 ethoxy)phenyl]methyl]amine product, as a light yellow oil, > 90% pure by HPLC analysis. HRMS calcd. for $C_{21}H_{15}NO_2BrF_5$: 488.0285 $[M+H]^+$, found: 488.0269.

- The 3-[3-(2-bromo-5-fluorophenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-
- 20 phenyl]-methyl]amine (0.5 g, 2.0 mmol) product from **EX-43B** and *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.17 g, 2.0 mmol) from **EX-4** were dissolved in 0.5 mL of acetonitrile. Ytterbium (III) trifluoromethanesulfonate (0.06 g, 0.1 mmol) was added, and the stirred solution was warmed to 40 °C for 1 h, at which time HPLC analysis indicated that no secondary amine starting material remained. The
- 25 reaction was quenched with water and extracted with ether. The ether layer was washed with water and brine, then dried over $MgSO_4$. The crude product was

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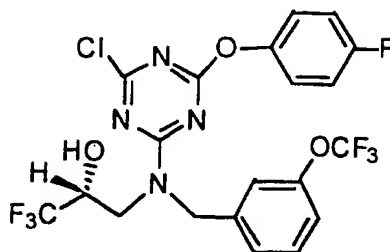
purified by flash column chromatography on silica gel eluting with 1:7:0.01 of ethyl acetate:hexane:ammonium hydroxide to give 0.4 g (67%) of the desired *R*-(+)-3-[[3-(2-bromo-5-fluorophenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl] amino]-1,1,1-trifluoro-2-propanol product as a light yellow oil (> 84% ee by chiral HPLC analysis). Anal. calcd. for $C_{24}H_{18}BrF_8NO_3$: C, 48.02; H, 3.02; N, 2.33. found: C, 48.07; H, 3.14; N, 2.31. HRMS calcd.: 600.0420 $[M+H]^+$, found: 600.0386. 1H NMR ($CDCl_3$) δ 7.5 0 (dd, 1H), 7.30 (t, 1H), 7.18 (t, 1H), 7.07 (t, 2H), 6.99 (s, 1H), 6.70 (dt, 1H), 6.56 (dd, 1H), 6.52 (dd, 1H), 6.38 (dd, 1H), 6.32 (m, 1H), 5.87 (tt, 1H), 4.65 (d, 2H), 4.33 (m, 1H), 3.85 (dd, 1H), 3.56 (dd, 1H), 2.48 (bs, 1H). NOE difference spectra confirmed that the isolated material was the indicated *N*-[3-(2-bromo-5-fluorophenoxy)phenyl]-3-aminopropanol product. ^{19}F NMR ($CDCl_3$) δ -79.24 (d, 3F), -88.57 (m, 2F), -112.04 (q, 1H), -137.16 (dt, 2F).

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EXAMPLE 44



5 **(2R)-N-[2-chloro-6-(p-fluorophenoxy)-1,3,5-triazin-4-yl]-3-[[[3-(trifluoromethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol**

EX-44A) 3-Trifluoromethoxybenzenemethanamine (1.15g, 6 mmol) and *R*-(+)-1,1,1-trifluoro-2,3-epoxypropane (0.67 g, 6 mmol) were combined and stirred at
 10 80 °C for 1.5 h. The mixture was cooled to room temperature, and the resulting solid was recrystallized from hot hexanes. The white solid was isolated by vacuum filtration and washed with cold hexanes to give 0.67 g (37%) of pure (2R)-3-[[[3-(trifluoro-methoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol.
¹H NMR (CDCl₃) δ 7.37 (t, 1H), 7.24 (d, 1H), 7.15 (m, 2H), 3.99 (m, 1H), 3.85
 15 (d, 2H), 2.98 (dd, 1H), 2.88 (dd, 1H), 2.79 (s, 1H). ¹⁹F NMR (CDCl₃) δ -58.19 (s, 3F), -78.88 (s, 3F). HRMS calcd. for C₁₁H₁₁F₆NO₂: 304.0772 [M+H]⁺, found: 304.0794.

EX-44B) To a solution of p-fluorophenol 1.00 g (8.92 mmol) in 30 mL of
 20 tetrahydrofuran at 0 °C was added a 60% dispersion of sodium hydride in mineral oil (0.36 g, 8.92 mmol). After 30 min, cyanuric chloride (1.64 g, 8.92 mmol) was added as a heterogeneous mixture in tetrahydrofuran at 0 °C. The reaction mixture was allowed to slowly warm to room temperature. After 14 h, the

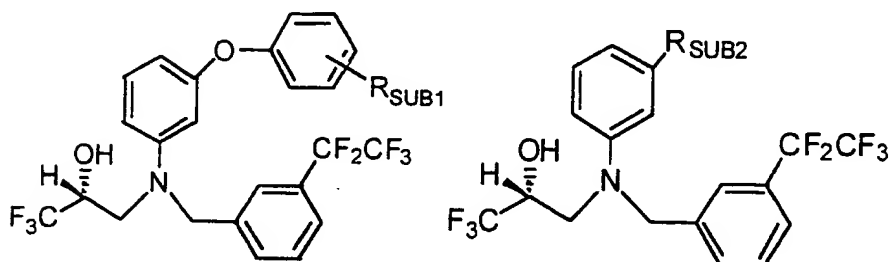
mixture was cooled to 0 °C. and a saturated aqueous NH₄Cl solution was added. The aqueous solution was extracted with diethyl ether (3 x 50 mL). The combined ether extracts were washed with brine, dried (MgSO₄), and concentrated *in vacuo* to afford 1.34 g (58%) of the desired 2,4-dichloro-6-(4-fluorophenoxy)-1,3,5-triazine product as an off white solid which was taken on to the next step without purification. MS *m/z* = 260 [M+H]⁺.

To a stirred solution of aminopropanol from EX-44A (0.100 g, 0.330 mmol) in *N,N*-dimethylformamide at 0 °C was added the 2,4-dichloro-(4-fluorophenoxy)-1,3,5-triazine ether product from EX-44B (0.086 g, 0.330 mmol) as a solution in *N,N*-di-methylformamide. The reaction mixture was allowed to slowly warm to room temperature. After 14 h, the reaction mixture was cooled to 0 °C, and a saturated aq. NaHCO₃ solution was added. After stirring the reaction mixture for 30 min at room temperature, the aqueous layer was extracted with ether (3 x 30 mL). The combined ether extracts were washed with brine, dried (MgSO₄), and concentrated *in vacuo* to give a yellow oil. The crude residue was purified by column chromatography on silica gel eluting with 20 % ethyl acetate in hexanes to give 0.075 g (43%) of the desired (2*R*)-*N*-[2-chloro-6-(*p*-fluorophenoxy)-1,3,5-triazin-4-yl]-3-[[[3-(trifluoromethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol product as a pale yellow oil. HRMS calcd. for C₂₀H₁₄ClF₇N₄O₃: 526.0643 [M⁺], found: 526.0632. ¹H NMR (C₆D₆) δ 6.95 (s, 1H), 6.63 (m, 14H), 4.74 (d, 1H), 4.37 (d, 1H), 4.16 (d, 1H), 4.00 (d, 2H), 3.73 (m, 1H), 3.48 (m, 2H), 3.26 (m, 2H), 3.12 (m, 2H).

Based on the preceding procedures, additional substituted (2*R*)-3-[(*N*-aryl)-[[aryl]methyl]amino]-1,1,1-trifluoro-2-propanols are prepared by one

skilled in the art using similar methods, as shown in Example Table 8. Substituted (3*R*)-4-[*N*-(aryl)-[(aryl)methyl]amino]-1.1.1.2.2-pentafluoro-3-butanols are prepared by one skilled in the art using similar methods, as shown in Example Table 9. Substituted (2*R*)-3-[*N*-(aryl)[(aryl)oxy]amino]-1.1.1-trifluoro-2-
5 propanols are prepared by one skilled in the art using similar methods, as shown in Example Table 10. Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl]amino]-1.1-difluoro-1-chloro-2-propanols are prepared by one skilled in the art using similar methods, as shown in Example Table 11. Substituted (2*R*)-3-[*N,N'*-(diaryl)amino]-
10 1.1.1-trifluoro-2-propanols are prepared by one skilled in the art using similar methods, as shown in Example Table 12.

Example Table 8. Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl]amino]-1,1,1-trifluoro-2-propanols.



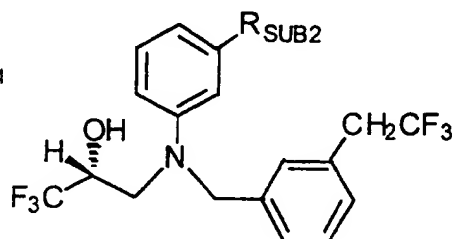
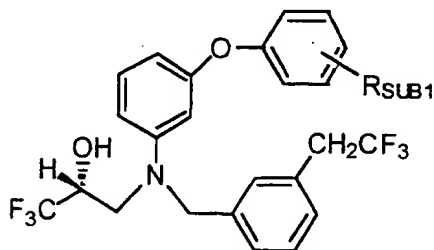
| Ex. No. | R _{SUB1} |
|---------|---|
| 45 | 3-isopropyl |
| 46 | 2-Cl, 3-Cl |
| 47 | 3-CF ₃ O |
| 48 | 4-F |
| 49 | 4-CH ₃ |
| 50 | 2-F, 5-Br |
| 51 | 3-CF ₃ CF ₂ |
| 52 | 3-CH ₃ CH ₂ |
| 53 | 3-CH ₃ , 5-CH ₃ |
| 54 | 3-(CH ₃) ₃ C |
| 55 | 4-F, 3-CH ₃ |
| 56 | 3-Cl, 4-Cl |
| 57 | 3,4-(CH ₂) ₄ |
| 58 | 3-HCF ₂ CF ₂ O |
| 59 | 3-CHF ₂ O |
| 60 | 3-(CH ₃) ₂ N |
| 61 | 3-cyclopropyl |
| 62 | 3-(2-furyl) |
| 63 | 3-CF ₃ CF ₂ |
| 64 | 4-NH ₂ |
| 65 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 69 | 3-CF ₃ O-benzyloxy |
| 70 | 3-CF ₃ -benzyloxy |
| 71 | 3-F, 5-F-benzyloxy |
| 72 | cyclohexylmethylenedioxy |
| 73 | benzyloxy |
| 74 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 75 | 4-CF ₃ O-benzyloxy |
| 76 | 4-CH ₃ CH ₂ -benzyloxy |
| 77 | isopropoxy |
| 78 | 3-CF ₃ -benzyl |
| 79 | isopropylthio |
| 80 | cyclopentoxo |
| 81 | 3-Cl-5-pyridinyloxy |
| 82 | 3-CF ₃ S-benzyloxy |
| 83 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 84 | 2-F, 3-CF ₃ -benzyloxy |
| 85 | 3-F, 5-CF ₃ -benzyloxy |
| 86 | 4-(CH ₃) ₂ CH-benzyloxy |
| 87 | 1-phenylethoxy |
| 88 | 4-F, 3-CH ₃ -benzoyl |
| 89 | 3-CF ₃ -phenyl |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 66 | 4-CH ₃ CH ₂ CH ₂ O |
| 67 | 3-CF ₃ |
| 68 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|---------------------------------|
| 90 | 4-CH ₃ O-phenylamino |
| 91 | cyclopropoxy |
| 92 | 4-NO ₂ -phenylthio |



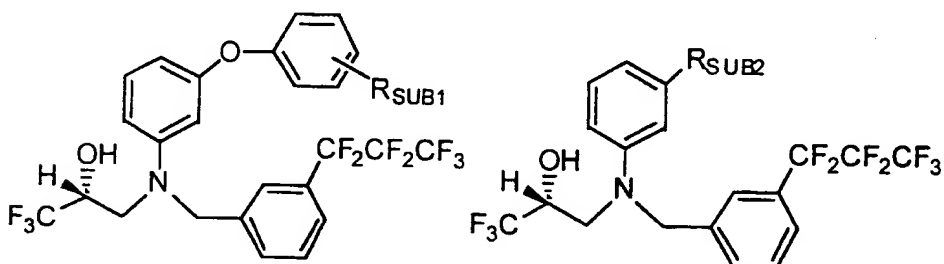
| Ex. No. | R _{SUB1} |
|---------|---|
| 93 | 3-isopropyl |
| 94 | 2-Cl, 3-Cl |
| 95 | 3-CF ₃ O |
| 96 | 4-F |
| 97 | 4-CH ₃ |
| 98 | 2-F, 5-Br |
| 99 | 4-Cl, 3-CH ₃ CH ₂ |
| 100 | 3-CH ₃ CH ₂ |
| 101 | 3-CH ₃ , 5-CH ₃ |
| 102 | 3-(CH ₃) ₃ C |
| 103 | 4-F, 3-CH ₃ |
| 104 | 3-Cl, 4-Cl |
| 105 | 3,4-(CH ₂) ₄ |
| 106 | 3-HCF ₂ CF ₂ O |
| 107 | 3-CHF ₂ O |

| Ex. No. | R _{SUB2} |
|---------|--|
| 117 | 3-CF ₃ O-benzyloxy |
| 118 | 3-CF ₃ -benzyloxy |
| 119 | 3-F, 5-F-benzyloxy |
| 120 | cyclohexylmethylenoxy |
| 121 | benzyloxy |
| 122 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 123 | 4-CF ₃ O-benzyloxy |
| 124 | 4-CH ₃ CH ₂ -benzyloxy |
| 125 | isopropoxy |
| 126 | 3-CF ₃ -benzyl |
| 127 | isopropylthio |
| 128 | cyclopentoxy |
| 129 | 3-Cl-5-pyridinyloxy |
| 130 | 3-CF ₃ S-benzyloxy |
| 131 | 3-CH ₃ , 4-CH ₃ -benzyloxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 108 | 3-(CH ₃) ₂ N |
| 109 | 3-cyclopropyl |
| 110 | 3-(2-furyl) |
| 111 | 3-CF ₃ CF ₂ |
| 112 | 4-NH ₂ |
| 113 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 114 | 4-CH ₃ CH ₂ CH ₂ O |
| 115 | 3-CF ₃ |
| 116 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 132 | 2-F, 3-CF ₃ -benzyloxy |
| 133 | 3-F, 5-CF ₃ -benzyloxy |
| 134 | 4-(CH ₃) ₂ CH-benzyloxy |
| 135 | 1-phenylethoxy |
| 136 | 4-F, 3-CH ₃ -benzoyl |
| 137 | 3-CF ₃ -phenyl |
| 138 | 4-CH ₃ O-phenylamino |
| 139 | cyclopropoxy |
| 140 | 4-NO ₂ -phenylthio |



| Ex. No. | R _{SUB1} |
|---------|---|
| 141 | 3-isopropyl |
| 142 | 2-Cl, 3-Cl |
| 143 | 3-CF ₃ O |
| 144 | 4-F |
| 145 | 4-CH ₃ |
| 146 | 2-F, 5-Br |
| 147 | 4-Cl, 3-CH ₃ CH ₂ |
| 148 | 3-CH ₃ CH ₂ |

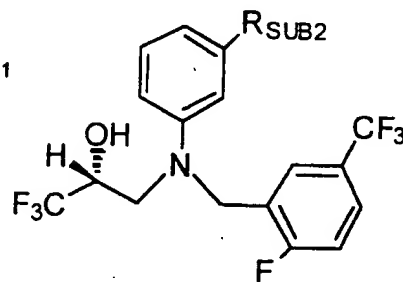
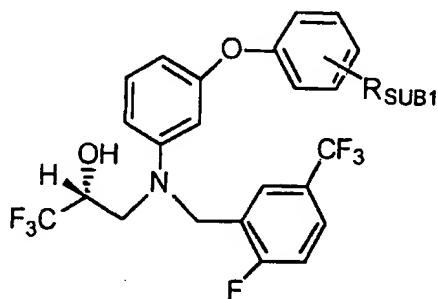
| Ex. No. | R _{SUB2} |
|---------|--|
| 165 | 3-CF ₃ O-benzyloxy |
| 166 | 3-CF ₃ -benzyloxy |
| 167 | 3-F, 5-F-benzyloxy |
| 168 | cyclohexylmethylenoxy |
| 169 | benzyloxy |
| 170 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 171 | 4-CF ₃ O-benzyloxy |
| 172 | 4-CH ₃ CH ₂ -benzyloxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 149 | 3-CH ₃ , 5-CH ₃ |
| 150 | 3-(CH ₃) ₃ C |
| 151 | 4-F, 3-CH ₃ |
| 152 | 3-Cl, 4-Cl |
| 153 | 3,4-(CH ₂) ₄ |
| 154 | 3-HCF ₂ CF ₂ O |
| 155 | 3-CHF ₂ O |
| 156 | 3-(CH ₃) ₂ N |
| 157 | 3-cyclopropyl |
| 158 | 3-(2-furyl) |
| 159 | 3-CF ₃ CF ₂ |
| 160 | 4-NH ₂ |
| 161 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 162 | 4-CH ₃ CH ₂ CH ₂ O |
| 163 | 3-CF ₃ |
| 164 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|--|
| 173 | isopropoxy |
| 174 | 3-CF ₃ -benzyl |
| 175 | isopropylthio |
| 176 | cyclopentoxy |
| 177 | 3-Cl-5-pyridinyloxy |
| 178 | 3-CF ₃ S-benzyloxy |
| 179 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 180 | 2-F, 3-CF ₃ -benzyloxy |
| 181 | 3-F, 5-CF ₃ -benzyloxy |
| 182 | 4-(CH ₃) ₂ CH-benzyloxy |
| 183 | 1-phenylethoxy |
| 184 | 4-F, 3-CH ₃ -benzoyl |
| 185 | 3-CF ₃ -phenyl |
| 186 | 4-CH ₃ O-phenylamino |
| 187 | cyclopropoxy |
| 188 | 4-NO ₂ -phenylthio |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.



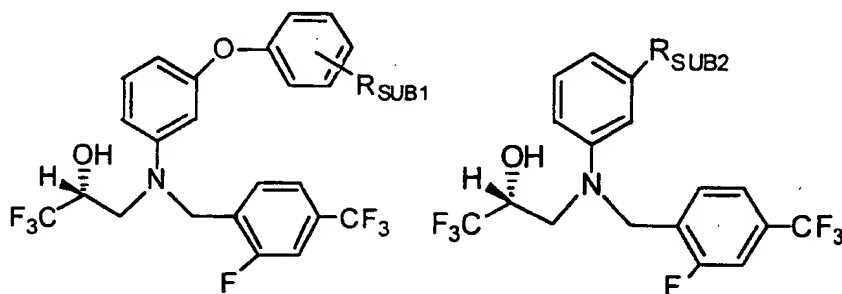
| Ex. No. | <u>R_{SUB1}</u> |
|---------|---------------------------------------|
| 189 | 3-isopropyl |
| 190 | 2-Cl, 3-Cl |
| 191 | 3-CF ₃ O |
| 192 | 4-F |
| 193 | 4-CH ₃ |
| 194 | 2-F, 5-Br |
| 195 | 4-Cl, 3-CH ₃ |
| 196 | 3-CH ₃ CH ₂ |
| 197 | 3-CH ₃ , 5-CH ₃ |
| 198 | 3-(CH ₃) ₃ C |
| 199 | 4-F, 3-CH ₃ |
| 200 | 3-Cl, 4-Cl |
| 201 | 3,4-(CH ₂) ₄ |
| 202 | 3-HCF ₂ CF ₂ O |
| 203 | 3-CHF ₂ O |
| 204 | 3-(CH ₃) ₂ N |
| 205 | 3-cyclopropyl |
| 206 | 3-(2-furyl) |
| 207 | 3-CF ₃ CF ₂ |

| Ex. No. | <u>R_{SUB2}</u> |
|---------|--|
| 213 | 3-CF ₃ O-benzyloxy |
| 214 | 3-CF ₃ -benzyloxy |
| 215 | 3-F, 5-F-benzyloxy |
| 216 | cyclohexylmethylenedioxy |
| 217 | benzyloxy |
| 218 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 219 | 4-CF ₃ O-benzyloxy |
| 220 | 4-CH ₃ CH ₂ -benzyloxy |
| 221 | isopropoxy |
| 222 | 3-CF ₃ -benzyl |
| 223 | isopropylthio |
| 224 | cyclopentoxy |
| 225 | 3-Cl-5-pyridinyloxy |
| 226 | 3-CF ₃ S-benzyloxy |
| 227 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 228 | 2-F, 3-CF ₃ -benzyloxy |
| 229 | 3-F, 5-CF ₃ -benzyloxy |
| 230 | 4-(CH ₃) ₂ CH-benzyloxy |
| 231 | 1-phenylethoxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 208 | 4-NH ₂ |
| 209 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 210 | 4-CH ₃ CH ₂ CH ₂ O |
| 211 | 3-CF ₃ |
| 212 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|---------------------------------|
| 232 | 4-F, 3-CH ₃ -benzoyl |
| 233 | 3-CF ₃ -phenyl |
| 234 | 4-CH ₃ O-phenylamino |
| 235 | cyclopropoxy |
| 236 | 4-NO ₂ -phenylthio |



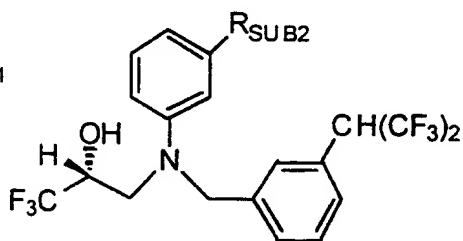
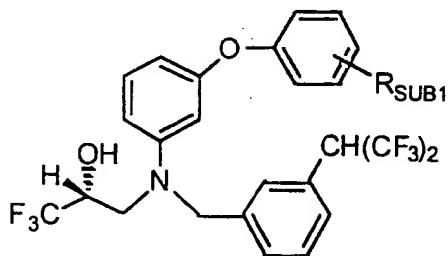
| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---------------------------------------|
| 237 | 3-isopropyl |
| 238 | 2-Cl, 3-Cl |
| 239 | 3-CF ₃ O |
| 240 | 4-F |
| 241 | 4-CH ₃ |
| 242 | 2-F, 5-Br |
| 243 | 4-Cl, 3-CH ₃ |
| 244 | 3-CH ₃ CH ₂ |
| 245 | 3-CH ₃ , 5-CH ₃ |
| 246 | 3-(CH ₃) ₃ C |
| 247 | 4-F, 3-CH ₃ |
| 248 | 3-Cl, 4-Cl |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 261 | 3-CF ₃ O-benzyloxy |
| 262 | 3-CF ₃ -benzyloxy |
| 263 | 3-F, 5-F-benzyloxy |
| 264 | cyclohexylmethylenedioxy |
| 265 | benzyloxy |
| 266 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 267 | 4-CF ₃ O-benzyloxy |
| 268 | 4-CH ₃ CH ₂ -benzyloxy |
| 269 | isopropoxy |
| 270 | 3-CF ₃ -benzyl |
| 271 | isopropylthio |
| 272 | cyclopentoxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 249 | 3,4-(CH ₂) ₄ |
| 250 | 3-HCF ₂ CF ₂ O |
| 251 | 3-CHF ₂ O |
| 252 | 3-(CH ₃) ₂ N |
| 253 | 3-cyclopropyl |
| 254 | 3-(2-furyl) |
| 255 | 3-CF ₃ CF ₂ |
| 256 | 4-NH ₂ |
| 257 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 258 | 4-CH ₃ CH ₂ CH ₂ O |
| 259 | 3-CF ₃ |
| 260 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 273 | 3-Cl-5-pyridinyloxy |
| 274 | 3-CF ₃ S-benzyloxy |
| 275 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 276 | 2-F, 3-CF ₃ -benzyloxy |
| 277 | 3-F, 5-CF ₃ -benzyloxy |
| 278 | 4-(CH ₃) ₂ CH-benzyloxy |
| 279 | 1-phenylethoxy |
| 280 | 4-F, 3-CH ₃ -benzoyl |
| 281 | 3-CF ₃ -phenyl |
| 282 | 4-CH ₃ O-phenylamino |
| 283 | cyclopropoxy |
| 284 | 4-NO ₂ -phenylthio |



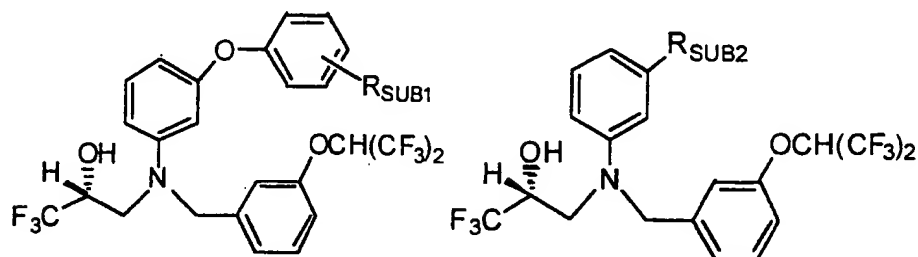
| Ex. No. | R _{SUB1} |
|---------|---------------------|
| 285 | 3-isopropyl |
| 286 | 2-Cl, 3-Cl |
| 287 | 3-CF ₃ O |
| 288 | 4-F |
| 289 | 4-CH ₃ |

| Ex. No. | R _{SUB2} |
|---------|-------------------------------|
| 309 | 3-CF ₃ O-benzyloxy |
| 310 | 3-CF ₃ -benzyloxy |
| 311 | 3-F, 5-F-benzyloxy |
| 312 | cyclohexylmethylenoxy |
| 313 | benzyloxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> | <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|---|--------------------|--|
| 290 | 2-F, 5-Br | 314 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 291 | 4-Cl, 3-CH ₃ CH ₂ | 315 | 4-CF ₃ O-benzyloxy |
| 292 | 3-CH ₃ CH ₂ | 316 | 4-CH ₃ CH ₂ -benzyloxy |
| 293 | 3-CH ₃ , 5-CH ₃ | 317 | isopropoxy |
| 294 | 3-(CH ₃) ₃ C | 318 | 3-CF ₃ -benzyl |
| 295 | 4-F, 3-CH ₃ | 319 | isopropylthio |
| 296 | 3-Cl, 4-Cl | 320 | cyclopentoxy |
| 297 | 3,4-(CH ₂) ₄ | 321 | 3-Cl-5-pyridinyloxy |
| 298 | 3-HCF ₂ CF ₂ O | 322 | 3-CF ₃ S-benzyloxy |
| 299 | 3-CHF ₂ O | 323 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 300 | 3-(CH ₃) ₂ N | 324 | 2-F, 3-CF ₃ -benzyloxy |
| 301 | 3-cyclopropyl | 325 | 3-F, 5-CF ₃ -benzyloxy |
| 302 | 3-(2-furyl) | 326 | 4-(CH ₃) ₂ CH-benzyloxy |
| 303 | 3-CF ₃ CF ₂ | 327 | 1-phenylethoxy |
| 304 | 4-NH ₂ | 328 | 4-F, 3-CH ₃ -benzoyl |
| 305 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ | 329 | 3-CF ₃ -phenyl |
| 306 | 4-CH ₃ CH ₂ CH ₂ O | 330 | 4-CH ₃ O-phenylamino |
| 307 | 3-CF ₃ | 331 | cyclopropoxy |
| 308 | 2-NO ₂ | 332 | 4-NO ₂ -phenylthio |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.



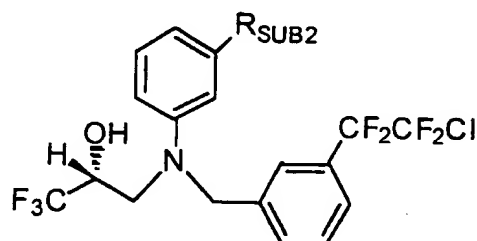
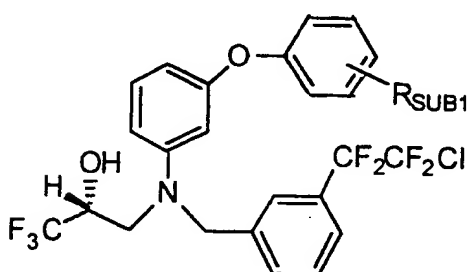
| Ex. No. | R _{SUB1} |
|---------|---|
| 333 | 3-isopropyl |
| 334 | 2-Cl, 3-Cl |
| 335 | 3-CF ₃ O |
| 336 | 4-F |
| 337 | 4-CH ₃ |
| 338 | 2-F, 5-Br |
| 339 | 4-Cl, 3-CH ₃ CH ₂ |
| 340 | 3-CH ₃ CH ₂ |
| 341 | 3-CH ₃ , 5-CH ₃ |
| 342 | 3-(CH ₃) ₃ C |
| 343 | 4-F, 3-CH ₃ |
| 344 | 3-Cl, 4-Cl |
| 345 | 3,4-(CH ₂) ₄ |
| 346 | 3-HCF ₂ CF ₂ O |
| 347 | 3-CHF ₂ O |
| 348 | 3-(CH ₃) ₂ N |
| 349 | 3-cyclopropyl |
| 350 | 3-(2-furyl) |
| 351 | 3-CF ₃ CF ₂ |
| 352 | 4-NH ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 357 | 3-CF ₃ O-benzyloxy |
| 358 | 3-CF ₃ -benzyloxy |
| 359 | 3-F, 5-F-benzyloxy |
| 360 | cyclohexylmethylenoxy |
| 361 | benzyloxy |
| 362 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 363 | 4-CF ₃ O-benzyloxy |
| 364 | 4-CH ₃ CH ₂ -benzyloxy |
| 365 | isopropoxy |
| 366 | 3-CF ₃ -benzyl |
| 367 | isopropylthio |
| 368 | cyclopentoxy |
| 369 | 3-Cl-5-pyridinyloxy |
| 370 | 3-CF ₃ S-benzyloxy |
| 371 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 372 | 2-F, 3-CF ₃ -benzyloxy |
| 373 | 3-F, 5-CF ₃ -benzyloxy |
| 374 | 4-(CH ₃) ₂ CH-benzyloxy |
| 375 | 1-phenylethoxy |
| 376 | 4-F, 3-CH ₃ -benzoyl |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 353 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 354 | 4-CH ₃ CH ₂ CH ₂ O |
| 355 | 3-CF ₃ |
| 356 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|---------------------------------|
| 377 | 3-CF ₃ -phenyl |
| 378 | 4-CH ₃ O-phenylamino |
| 379 | cyclopropoxy |
| 380 | 4-NO ₂ -phenylthio |



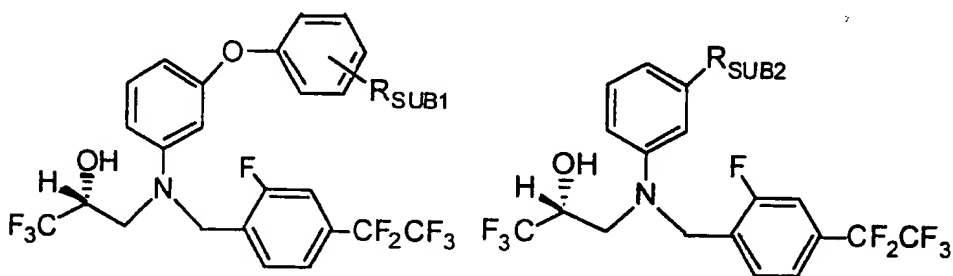
| Ex. No. | R _{SUB1} |
|---------|---|
| 381 | 3-isopropyl |
| 382 | 2-Cl, 3-Cl |
| 383 | 3-CF ₃ O |
| 384 | 4-F |
| 385 | 4-CH ₃ |
| 386 | 2-F, 5-Br |
| 387 | 4-Cl, 3-CH ₃ CH ₂ |
| 388 | 3-CH ₃ CH ₂ |
| 389 | 3-CH ₃ , 5-CH ₃ |
| 390 | 3-(CH ₃) ₃ C |
| 391 | 4-F, 3-CH ₃ |
| 392 | 3-Cl, 4-Cl |
| 393 | 3,4-(CH ₂) ₄ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 405 | 3-CF ₃ O-benzyloxy |
| 406 | 3-CF ₃ -benzyloxy |
| 407 | 3-F, 5-F-benzyloxy |
| 408 | cyclohexylmethylenoxy |
| 409 | benzyloxy |
| 410 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 411 | 4-CF ₃ O-benzyloxy |
| 412 | 4-CH ₃ CH ₂ -benzyloxy |
| 413 | isopropoxy |
| 414 | 3-CF ₃ -benzyl |
| 415 | isopropylthio |
| 416 | cyclopentoxy |
| 417 | 3-Cl-5-pyridinyloxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 394 | 3-HCF ₂ CF ₂ O |
| 395 | 3-CHF ₂ O |
| 396 | 3-(CH ₃) ₂ N |
| 397 | 3-cyclopropyl |
| 398 | 3-(2-furyl) |
| 399 | 3-CF ₃ CF ₂ |
| 400 | 4-NH ₂ |
| 401 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 402 | 4-CH ₃ CH ₂ CH ₂ O |
| 403 | 3-CF ₃ |
| 404 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 418 | 3-CF ₃ S-benzyloxy |
| 419 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 420 | 2-F, 3-CF ₃ -benzyloxy |
| 421 | 3-F, 5-CF ₃ -benzyloxy |
| 422 | 4-(CH ₃) ₂ CH-benzyloxy |
| 423 | 1-phenylethoxy |
| 424 | 4-F, 3-CH ₃ -benzoyl |
| 425 | 3-CF ₃ -phenyl |
| 426 | 4-CH ₃ O-phenylamino |
| 427 | cyclopropoxy |
| 428 | 4-NO ₂ -phenylthio |



| Ex. No. | R _{SUB1} |
|---------|---------------------|
| 429 | 3-isopropyl |
| 430 | 2-Cl, 3-Cl |
| 431 | 3-CF ₃ O |
| 432 | 4-F |
| 433 | 4-CH ₃ |
| 434 | 2-F, 5-Br |

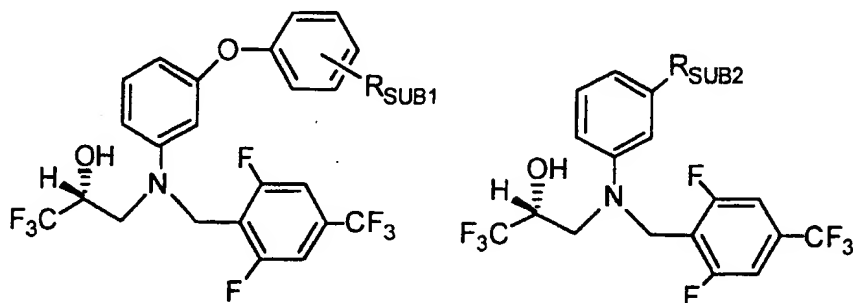
| Ex. No. | R _{SUB2} |
|---------|--|
| 453 | 3-CF ₃ O-benzyloxy |
| 454 | 3-CF ₃ -benzyloxy |
| 455 | 3-F, 5-F-benzyloxy |
| 456 | cyclohexylmethyleneoxy |
| 457 | benzyloxy |
| 458 | 3-CF ₃ , 5-CF ₃ -benzyloxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 435 | 4-Cl, 3-CH ₃ CH ₂ |
| 436 | 3-CH ₃ CH ₂ |
| 437 | 3-CH ₃ , 5-CH ₃ |
| 438 | 3-(CH ₃) ₃ C |
| 439 | 4-F, 3-CH ₃ |
| 440 | 3-Cl, 4-Cl |
| 441 | 3,4-(CH ₂) ₄ |
| 442 | 3-HCF ₂ CF ₂ O |
| 443 | 3-CHF ₂ O |
| 444 | 3-(CH ₃) ₂ N |
| 445 | 3-cyclopropyl |
| 446 | 3-(2-furyl) |
| 447 | 3-CF ₃ CF ₂ |
| 448 | 4-NH ₂ |
| 449 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 450 | 4-CH ₃ CH ₂ CH ₂ O |
| 451 | 3-CF ₃ |
| 452 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|--|
| 459 | 4-CF ₃ O-benzyloxy |
| 460 | 4-CH ₃ CH ₂ -benzyloxy |
| 461 | isopropoxy |
| 462 | 3-CF ₃ -benzyl |
| 463 | isopropylthio |
| 464 | cyclopentoxy |
| 465 | 3-Cl-5-pyridinyloxy |
| 466 | 3-CF ₃ S-benzyloxy |
| 467 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 468 | 2-F, 3-CF ₃ -benzyloxy |
| 469 | 3-F, 5-CF ₃ -benzyloxy |
| 470 | 4-(CH ₃) ₂ CH-benzyloxy |
| 471 | 1-phenylethoxy |
| 472 | 4-F, 3-CH ₃ -benzoyl |
| 473 | 3-CF ₃ -phenyl |
| 474 | 4-CH ₃ O-phenylamino |
| 475 | cyclopropoxy |
| 476 | 4-NO ₂ -phenylthio |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-{(aryl)methyl} amino]-1,1,1-trifluoro-2-propanols.



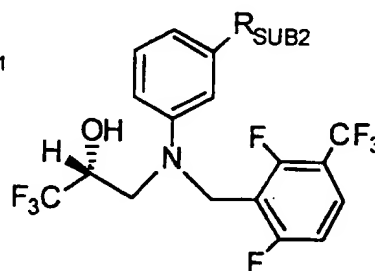
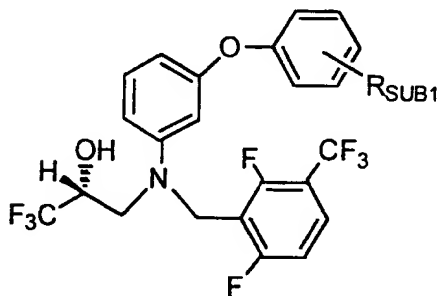
| Ex. No. | R _{SUB1} |
|---------|---|
| 477 | 3-isopropyl |
| 478 | 2-Cl, 3-Cl |
| 479 | 3-CF ₃ O |
| 480 | 4-F |
| 481 | 4-CH ₃ |
| 482 | 2-F, 5-Br |
| 483 | 4-Cl, 3-CH ₃ CH ₂ |
| 484 | 3-CH ₃ CH ₂ |
| 485 | 3-CH ₃ , 5-CH ₃ |
| 486 | 3-(CH ₃) ₃ C |
| 487 | 4-F, 3-CH ₃ |
| 488 | 3-Cl, 4-Cl |
| 489 | 3,4-(CH ₂) ₄ |
| 490 | 3-HCF ₂ CF ₂ O |
| 491 | 3-CHF ₂ O |
| 492 | 3-(CH ₃) ₂ N |
| 493 | 3-cyclopropyl |
| 494 | 3-(2-furyl) |
| 495 | 3-CF ₃ CF ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 501 | 3-CF ₃ O-benzyloxy |
| 502 | 3-CF ₃ -benzyloxy |
| 503 | 3-F, 5-F-benzyloxy |
| 504 | cyclohexylmethylenedioxy |
| 505 | benzyloxy |
| 506 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 507 | 4-CF ₃ O-benzyloxy |
| 508 | 4-CH ₃ CH ₂ -benzyloxy |
| 509 | isopropoxy |
| 510 | 3-CF ₃ -benzyl |
| 511 | isopropylthio |
| 512 | cyclopentoxy |
| 513 | 3-Cl-5-pyridinyloxy |
| 514 | 3-CF ₃ S-benzyloxy |
| 515 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 516 | 2-F, 3-CF ₃ -benzyloxy |
| 517 | 3-F, 5-CF ₃ -benzyloxy |
| 518 | 4-(CH ₃) ₂ CH-benzyloxy |
| 519 | 1-phenylethoxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-{(aryl)methyl} amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 496 | 4-NH ₂ |
| 497 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 498 | 4-CH ₃ CH ₂ CH ₂ O |
| 499 | 3-CF ₃ |
| 500 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|---------------------------------|
| 520 | 4-F, 3-CH ₃ -benzoyl |
| 521 | 3-CF ₃ -phenyl |
| 522 | 4-CH ₃ O-phenylamino |
| 523 | cyclopropoxy |
| 524 | 4-NO ₂ -phenylthio |



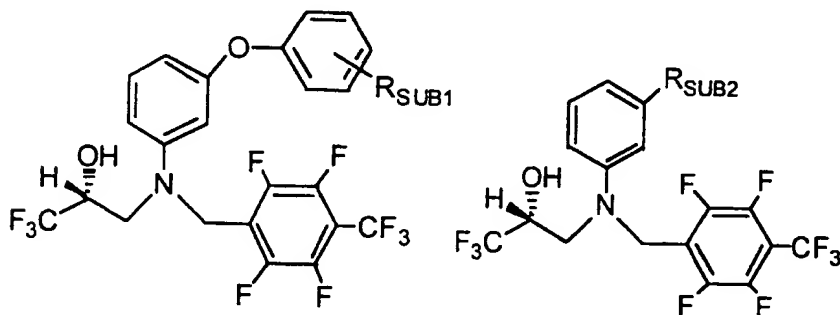
| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 525 | 3-isopropyl |
| 526 | 2-Cl, 3-Cl |
| 527 | 3-CF ₃ O |
| 528 | 4-F |
| 529 | 4-CH ₃ |
| 530 | 2-F, 5-Br |
| 531 | 4-Cl, 3-CH ₃ CH ₂ |
| 532 | 3-CH ₃ CH ₂ |
| 533 | 3-CH ₃ , 5-CH ₃ |
| 534 | 3-(CH ₃) ₃ C |
| 535 | 4-F, 3-CH ₃ |
| 536 | 3-Cl, 4-Cl |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 549 | 3-CF ₃ O-benzyloxy |
| 550 | 3-CF ₃ -benzyloxy |
| 551 | 3-F, 5-F-benzyloxy |
| 552 | cyclohexylmethylenoxy |
| 553 | benzyloxy |
| 554 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 555 | 4-CF ₃ O-benzyloxy |
| 556 | 4-CH ₃ CH ₂ -benzyloxy |
| 557 | isopropoxy |
| 558 | 3-CF ₃ -benzyl |
| 559 | isopropylthio |
| 560 | cyclopentoxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 537 | 3,4-(CH ₂) ₄ |
| 538 | 3-HCF ₂ CF ₂ O |
| 539 | 3-CHF ₂ O |
| 540 | 3-(CH ₃) ₂ N |
| 541 | 3-cyclopropyl |
| 542 | 3-(2-furyl) |
| 543 | 3-CF ₃ CF ₂ |
| 544 | 4-NH ₂ |
| 545 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 546 | 4-CH ₃ CH ₂ CH ₂ O |
| 547 | 3-CF ₃ |
| 548 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 561 | 3-Cl-5-pyridinyloxy |
| 562 | 3-CF ₃ S-benzyloxy |
| 563 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 564 | 2-F, 3-CF ₃ -benzyloxy |
| 565 | 3-F, 5-CF ₃ -benzyloxy |
| 566 | 4-(CH ₃) ₂ CH-benzyloxy |
| 567 | 1-phenylethoxy |
| 568 | 4-F, 3-CH ₃ -benzoyl |
| 569 | 3-CF ₃ -phenyl |
| 570 | 4-CH ₃ O-phenylamino |
| 571 | cyclopropoxy |
| 572 | 4-NO ₂ -phenylthio |



| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|-------------------------|
| 573 | 3-isopropyl |
| 574 | 2-Cl, 3-Cl |
| 575 | 3-CF ₃ O |
| 576 | 4-F |

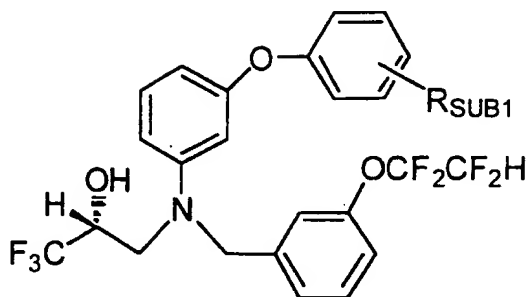
| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|-------------------------------|
| 597 | 3-CF ₃ O-benzyloxy |
| 598 | 3-CF ₃ -benzyloxy |
| 599 | 3-F, 5-F-benzyloxy |
| 600 | cyclohexylmethylenoxy |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 577 | 4-CH ₃ |
| 578 | 2-F, 5-Br |
| 579 | 4-Cl, 3-CH ₃ CH ₂ |
| 580 | 3-CH ₃ CH ₂ |
| 581 | 3-CH ₃ , 5-CH ₃ |
| 582 | 3-(CH ₃) ₃ C |
| 583 | 4-F, 3-CH ₃ |
| 584 | 3-Cl, 4-Cl |
| 585 | 3,4-(CH ₂) ₄ |
| 586 | 3-HCF ₂ CF ₂ O |
| 587 | 3-CHF ₂ O |
| 588 | 3-(CH ₃) ₂ N |
| 589 | 3-cyclopropyl |
| 590 | 3-(2-furyl) |
| 591 | 3-CF ₃ CF ₂ |
| 592 | 4-NH ₂ |
| 593 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 594 | 4-CH ₃ CH ₂ CH ₂ O |
| 595 | 3-CF ₃ |
| 596 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|--|
| 601 | benzyloxy |
| 602 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 603 | 4-CF ₃ O-benzyloxy |
| 604 | 4-CH ₃ CH ₂ -benzyloxy |
| 605 | isopropoxy |
| 606 | 3-CF ₃ -benzyl |
| 607 | isopropylthio |
| 608 | cyclopentoxy |
| 609 | 3-Cl-5-pyridinyloxy |
| 610 | 3-CF ₃ S-benzyloxy |
| 611 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 612 | 2-F, 3-CF ₃ -benzyloxy |
| 613 | 3-F, 5-CF ₃ -benzyloxy |
| 614 | 4-(CH ₃) ₂ CH-benzyloxy |
| 615 | 1-phenylethoxy |
| 616 | 4-F, 3-CH ₃ -benzoyl |
| 617 | 3-CF ₃ -phenyl |
| 618 | 4-CH ₃ O-phenylamino |
| 619 | cyclopropoxy |
| 620 | 4-NO ₂ -phenylthio |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.



| <u>Ex. No.</u> | <u>R_{SUB1}</u> | <u>Calculated Mass [M+H]⁺</u> | <u>Observed Mass [M+H]⁺</u> |
|--------------------|---------------------------------------|--|--|
| 621 | 4-F | 522.1315 | 522.1297 |
| 622 | 2-Cl, 3-Cl | 572.0630 | 572.0653 |
| 623 | 2-F, 5-Br | 600.0420 | 600.0404 |
| 624 | 4-Cl, 3-CH ₃ | 551.1098 | 551.1101 |
| 625 | 3-CH ₃ , 5-CH ₃ | 532.1722 | 532.1705 |
| 626 | 3-(CH ₃) ₃ C | 560.2035 | 560.2055 |
| 627 | 4-F, 3-CH ₃ | 536.1471 | 536.1480 |
| 628 | 3-Cl, 4-Cl | 572.0630 | 572.0630 |
| 629 | 3,4-(CH ₂) ₄ | 558.1879 | 558.1881 |
| 630 | 3-HCF ₂ CF ₂ O | | |
| 631 | 3-CHF ₂ O | | |
| 632 | 3-(CH ₃) ₂ N | 547.1831 | 547.1844 |
| 633 | 3-cyclopropyl | | |
| 634 | 3-(2-furyl) | | |
| 635 | 3-CF ₃ CF ₂ | | |
| 636 | 3-cyclopentyl | | |
| 637 | 4-NH ₂ | 519.1519 | 519.1529 |

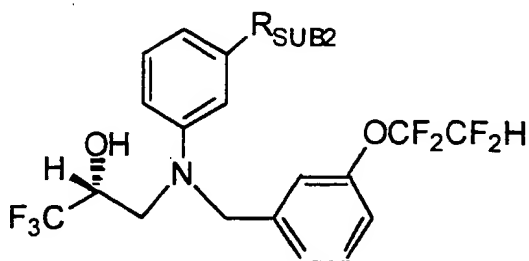
Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-{(aryl)methyl} amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> | <u>Calculated Mass [M+H]⁺</u> | <u>Observed Mass [M+H]⁺</u> |
|--------------------|---|--|--|
| 638 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ | 546.1879 | 546.1901 |
| 639 | 4-CH ₃ CH ₂ O | 547.1594 | 547.1594 |
| 640 | 3-CF ₃ | | |
| 641 | 2-NO ₂ | 549.1260 | 549.1235 |
| 642 | 3,4-dimethyl | 531.1644 | 531.1649 |
| 643 | 3-methyl, 5-ethyl | 546.1879 | 546.1899 |
| 644 | 3-methyl | 517.1488 | 517.1493 |
| 645 | 2,3-difluoro | 540.1221 | 540.1182 |
| 646 | 4-CF ₃ | 572.1282 | 572.1268 |
| 647 | 2-fluoro, 3-CF ₃ | 590.1189 | 590.1184 |
| 648 | 2-fluoro, 4-CF ₃ | 590.1189 | 590.1155 |
| 649 | 2-chloro, 4-fluoro | 556.0925 | 556.0891 |
| 650 | 4- <i>n</i> -propyl | 546.1879 | 546.1878 |
| 651 | 3-chloro, 4-fluoro | 556.0925 | 556.0932 |
| 652 | 2,4-difluoro | 540.1221 | 540.1194 |
| 653 | 3,5-difluoro | 540.1221 | 540.1217 |
| 654 | 3,4-difluoro | 540.1221 | 540.1248 |
| 655 | 3-fluoro | 522.1315 | 522.1337 |
| 656 | 2-chloro | 538.1019 | 538.1021 |
| 657 | 2-fluoro | 522.1315 | 522.1310 |
| 658 | 2,5-difluoro | 540.1221 | 540.1255 |
| 659 | 4-chloro, 2-fluoro | 556.0926 | 556.0954 |
| 660 | 2,4-dichloro | 572.0630 | 572.0667 |
| 661 | 2-fluoro, 3-CH ₃ | | |
| 662 | 4-chloro | 537.0942 | 537.0944 |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex.</u> <u>No.</u> | <u>R_{SUB1}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|---------------------------------------|--|--|
| 663 | 4-isopropyl, 3-methyl | 560.2035 | 560.2035 |
| 664 | 2,3,4-trifluoro | 558.1127 | 558.1161 |
| 665 | 2,3,5-trifluoro | 558.1127 | 558.1109 |
| 666 | 4-propoxy | 562.1828 | 562.1803 |
| 667 | 4-isopropyl | 546.1879 | 546.1899 |
| 668 | 4-CF ₃ O- | 588.1233 | 588.1241 |
| 669 | 4-butoxy | 576.1958 | 576.1969 |
| 670 | 3-methyl, 4-CH ₃ S- | 564.1443 | 564.1476 |
| 671 | 4-nitro | 549.1260 | 549.1306 |
| 672 | 3-CF ₃ S- | | |
| 673 | 4-chloro, 3-fluoro | 556.0925 | 556.0933 |
| 674 | 3,5-dimethoxy | 564.1623 | 564.1617 |
| 675 | 4-bromo | 582.0716 | 582.0473 |
| 676 | 4- <i>sec</i> -butyl | 560.2035 | 560.2051 |
| 677 | 3-fluoro-2-nitro | 567.1166 | 567.1135 |
| 678 | 3-methoxy | 533.1437 | 533.1450 |
| 679 | 4-bromo-2-nitro | 627.0366 | 627.0375 |
| 680 | 4-cyano | 529.1362 | 529.1364 |
| 681 | 4-CH ₃ S- | 550.1209 | 550.1251 |
| 682 | 3,4-(CH=CH) ₂ | 554.1566 | 554.1578 |
| 683 | 4-CH ₃ CH ₂ NH- | 547.1832 | 547.1819 |
| 684 | 4-propionyl | 560.1672 | 560.1694 |
| 685 | 3-phenyl | 580.1723 | 580.1772 |
| 686 | 4-cyclopentyl | 572.2035 | 572.2029 |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.



| <u>Ex.</u> <u>No.</u> | <u>R_{SUB}2</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|-----------------------------------|--|--|
| 687 | 6-methyl-3-pyridinyloxy | 518.1440 | 518.1452 |
| 688 | 5-chloro-3-pyridinyloxy | 539.0972 | 539.1002 |
| 689 | 3-pyridinyloxy | 505.1362 | 505.1369 |
| 690 | 2-methyl-3-pyridinyloxy | 519.1518 | 519.1517 |
| 691 | 5-indolinylloxy | 543.1519 | 543.1630 |
| 692 | 4-fluoro-2-pyridinyloxy | 523.1268 | 523.1243 |
| 693 | 2-cyano-3-pyridinyloxy | 530.1315 | 530.1300 |
| 694 | 5-bromo-2-pyridinyloxy | 583.0667 | 583.0405 |
| 695 | 3-CF ₃ -2-pyridinyloxy | 573.1236 | 573.1205 |
| 696 | 2-pyridinylmethylenoxy | 519.1519 | 519.1522 |
| 697 | cyclohexylmethylenoxy | 524.2036 | 524.2028 |
| 698 | isopropoxy | 470.1488 | 470.1565 |
| 699 | cyclopentyloxy | 496.1723 | 496.1719 |
| 700 | neo-pentoxy | 498.1879 | 498.1845 |
| 701 | 4-(methoxycarbonyl)-butoxy | 542.1777 | 542.1827 |
| 702 | trifluoromethoxy | 496.0971 | 496.0959 |
| 703 | 2-methylpropoxy | 484.1723 | 484.1718 |
| 704 | 2-methoxyethoxy | 486.1515 | 486.1537 |
| 705 | 2-oxobutoxy | 498.1515 | 498.1529 |
| 706 | cyclohexyloxy | 510.1880 | 510.1910 |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex.</u> <u>No.</u> | <u>R_{SUB2}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|--|--|--|
| 707 | (methoxycarbonyl)methoxy | 500.1308 | 500.1297 |
| 708 | 4-tetrahydropyranyloxy | 512.1672 | 512.1631 |
| 709 | 1-phenylethoxy | 532.1723 | 532.1711 |
| 710 | 3-CF ₃ O-benzyloxy | 602.1389 | 602.1380 |
| 711 | 3-trifluoromethyl-benzyloxy | 586.1440 | 586.1419 |
| 712 | 3,5-dimethyl-benzyloxy | 546.1879 | 546.1890 |
| 713 | 3-bromo-benzyloxy | 596.0671 | 596.0641 |
| 714 | 3-CF ₃ S-benzyloxy | 618.1161 | 618.1151 |
| 715 | 3,4-dimethyl-benzyloxy | 546.1879 | 546.1881 |
| 716 | 3,5-difluoro-benzyloxy | 554.1378 | 554.1390 |
| 717 | 2-fluoro-3-CF ₃ -benzyloxy | 604.1346 | 604.1329 |
| 718 | benzyloxy | 518.1566 | 518.1578 |
| 719 | 3,5-(CF ₃) ₂ -benzyloxy | 654.1314 | 654.1308 |
| 720 | 3-fluoro-5-CF ₃ -benzyloxy | 604.1346 | 604.1309 |
| 721 | 4-CF ₃ O-benzyloxy | 602.1389 | 602.1383 |
| 722 | 3-chloro-benzyloxy | 552.1176 | 552.1157 |
| 723 | 4-ethyl-benzyloxy | 546.1879 | 546.1862 |
| 724 | 3-methyl-benzyloxy | 532.1723 | 532.1692 |
| 725 | 2-fluoro-benzyloxy | 536.1472 | 536.1465 |
| 726 | 2,3-difluoro-benzyloxy | 554.1378 | 554.1364 |
| 727 | 4-isopropyl-benzyloxy | 560.2036 | 560.2020 |
| 728 | 4-methyl-benzyloxy | 532.1723 | 532.1729 |
| 729 | 4-bromo-benzyloxy | 596.0671 | 596.0669 |
| 730 | 4-CF ₃ -benzyloxy | 586.1440 | 586.1400 |
| 731 | 4-fluoro-benzyloxy | 536.1472 | 536.1454 |

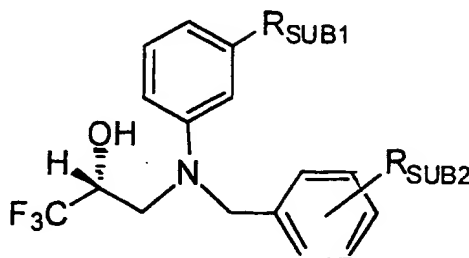
Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB2}</u> | <u>Calculated Mass [M+H]⁺</u> | <u>Observed Mass [M+H]⁺</u> |
|--------------------|--|--|--|
| 732 | 3-iodo-benzyloxy | 644.0533 | 644.0517 |
| 733 | 4-CF ₃ S-benzyloxy | 618.1161 | 618.1165 |
| 734 | 4-CF ₂ HO-benzyloxy | 584.1483 | 584.1480 |
| 735 | 4-fluoro-3-CF ₃ -benzyloxy | 604.1346 | 604.1336 |
| 736 | 2,3,5-trifluoro-benzyloxy | 572.1284 | 572.1276 |
| 737 | 4-chloro-benzyloxy | 552.1176 | 552.1188 |
| 738 | 2,5-difluoro-benzyloxy | 554.1378 | 554.1350 |
| 739 | 3-chloro-2-fluoro-benzyloxy | 570.1082 | 570.1069 |
| 740 | 2,4-(CF ₃) ₂ -benzyloxy | 654.1314 | 654.1321 |
| 741 | 3,5-dichloro-benzyloxy | 586.1787 | 586.1378 |
| 742 | 3-methoxy-benzyloxy | 548.1672 | 548.1676 |
| 743 | 4-cyano-benzyloxy | 543.1519 | 543.1517 |
| 744 | 4- <i>tert</i> -butyl-benzyloxy | 574.2192 | 574.2163 |
| 745 | isopropylthio | 486.1338 | 486.1351 |
| 746 | 4-nitrophenylthio | 565.1032 | 565.1034 |
| 747 | 4-acetylphenylthio | 562.1287 | 562.1261 |
| 748 | (4-chloro-thien-2-yl)- methylthio | 574.0512 | 574.0523 |
| 749 | 4-methoxy-phenylamino | 532.1597 | 532.1592 |
| 750 | 3-methoxy-phenylamino | 532.1597 | 532.1593 |
| 751 | 4-chloro-phenylamino | 536.1102 | 536.1125 |
| 752 | 4- <i>n</i> -propyl-phenylamino | 544.1961 | 544.1959 |
| 753 | 3-cyano-phenylamino | 527.1444 | 527.1448 |
| 754 | 3-CF ₃ -benzyl | 570.1413 | 570.1480 |
| 755 | 3-methyl-4-fluoro-benzyl | 534.1679 | 534.1688 |
| 756 | 3-CF ₃ -phenyl | 556.1334 | 556.1339 |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.

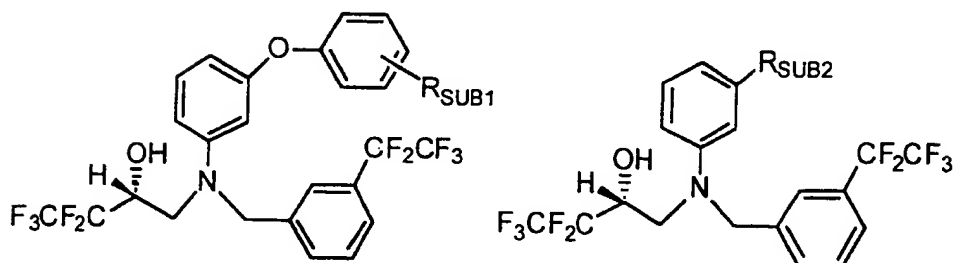
| <u>Ex.</u> <u>No.</u> | <u>R_{SUB2}</u> | <u>Calculated</u> <u>Mass</u> <u>[M+H]⁺</u> | <u>Observed</u> <u>Mass</u> <u>[M+H]⁺</u> |
|--------------------------|---------------------------|--|--|
| 757 | 2,4-dichloro-phenyl | 556.0681 | 556.0651 |
| 758 | 3-methoxybenzyl | 532.1723 | 532.1705 |
| 759 | 4-methoxyphenyl | 518.1566 | 518.1533 |
| 760 | 3-chloro-4-fluoro-phenyl | 540.0976 | 540.0957 |
| 761 | 4-fluoro-3-methyl-benzoyl | 548.1410 | 548.1441 |
| 762 | 3-chlorobenzyl | 536.1227 | 536.1218 |
| 763 | 3,4-dimethylbenzyl | 530.1930 | 530.1887 |
| 764 | 3,5-dichlorobenzyl | 570.0838 | 570.0801 |
| 765 | 2,3,4-trifluorophenyl | 542.1177 | 542.1152 |
| 766 | 3-chloro-4-fluoro-benzyl | 554.1133 | 554.1108 |
| 767 | 4-fluoro-3-methyl-phenyl | 520.1523 | 520.1494 |
| 768 | 3-methyl-4-chloro-benzyl | 550.1384 | 550.1380 |
| 769 | 2-methylpropanoyl | 482.1566 | 482.1576 |
| 770 | 4-methylthiobenzyl | 548.1494 | 548.1503 |
| 771 | 4-fluorophenyl | 506.1366 | 506.1336 |
| 772 | 4-chlorophenyl | 522.1071 | 522.1049 |
| 773 | 3-methoxyphenyl | 518.1566 | 518.1544 |
| 774 | 4-methylbenzyl | 516.1774 | 516.1769 |
| 775 | 1-hydroxy-2-methyl-propyl | 484.1723 | 484.1725 |
| 776 | benzyl | 502.1617 | 502.1609 |
| 777 | 2-CF ₃ -phenyl | 556.1334 | 556.1286 |
| 778 | 3,4-dichlorophenyl | 556.0681 | 556.0698 |
| 779 | benzoyl | 516.1410 | 516.1383 |
| 780 | 4-fluorobenzoyl | 534.1315 | 534.1273 |
| 781 | N-piperidinyl | 494.1804 | 494.1804 |
| 782 | phenyl | 488.1460 | 488.1457 |
| 783 | thien-2-yl | 494.1024 | 494.0987 |

Example Table 8 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)methyl] amino]-1,1,1-trifluoro-2-propanols.



| <u>Ex. No.</u> | <u>R_{SUB1}</u> | <u>R_{SUB2}</u> | <u>Calculated Mass [M+H]⁺</u> | <u>Observed Mass [M+H]⁺</u> |
|--------------------|--------------------------------|--|--|--|
| 784 | phenoxy | 3-cyclopentyl | 456.2150 | 456.2143 |
| 785 | phenoxy | 3-isopropoxy | 446.1943 | 446.1936 |
| 786 | phenoxy | 3-CF ₃ S | 488.1119 | 488.1116 |
| 787 | 4-F-phenoxy | 3-CF ₃ S | 505.0946 | 505.0927 |
| 788 | 4-F-phenoxy | 3- <i>sec</i> -butoxy | 478.2005 | 478.1880 |
| 789 | phenoxy | 3-(CF ₃) ₂ COH- | 554.1378 | 554.1385 |
| 790 | 4-CH ₃ - phenoxy | 3-CF ₃ S | 502.1275 | 502.1261 |
| 791 | phenoxy | 3-(2-furyl) | 454.1630 | 454.1635 |
| 792 | 4-F-phenoxy | 3-isopropoxy | 464.1849 | 464.1867 |
| 793 | phenoxy | 3-isobutyl | 444.2150 | 444.2157 |
| 794 | phenoxy | 3- <i>tert</i> -butoxy | 460.2100 | 460.2103 |
| 795 | 4-F-phenoxy | 3-CH ₃ CH ₂ O- | 450.1692 | 450.1682 |
| 796 | 4-F-phenoxy | 3-CF ₃ O- | 490.1253 | 490.1211 |
| 797 | phenoxy | 4-F-3-(2-furyl)- | 472.1536 | 472.1530 |
| 798 | 4-F-phenoxy | 3- <i>n</i> -propoxy- | 464.1849 | 464.1820 |
| 799 | 4-F-phenoxy | 3-cyclopentyloxy- | 490.2005 | 490.1998 |
| 800 | phenoxy | 3-(3-furyl)- | 454.1630 | 454.1646 |
| 801 | 4-F-phenoxy | 3-cyclopropyl- methyleneoxy | 476.1849 | 476.1857 |
| 802 | phenoxy | 3-CF ₃ CH ₂ O- | 486.1504 | 486.1498 |

Example Table 9. (3*R*)-4-[*N*-(aryl)-[(aryl)methyl]amino]-
1,1,1,2,2-pentafluoro-3-butanols.



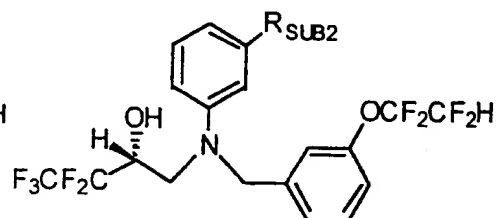
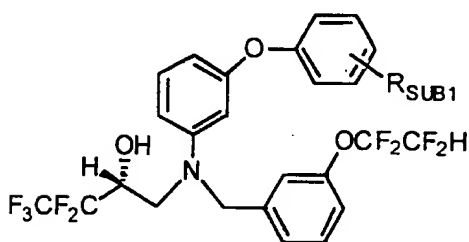
| Ex. No. | R _{SUB1} |
|---------|---|
| 803 | 3-isopropyl |
| 804 | 2-Cl, 3-Cl |
| 805 | 3-CF ₃ O |
| 806 | 4-F |
| 807 | 4-CH ₃ |
| 808 | 2-F, 5-Br |
| 809 | 4-Cl, 3-CH ₃ CH ₂ |
| 810 | 3-CH ₃ CH ₂ |
| 811 | 3-CH ₃ , 5-CH ₃ |
| 812 | 3-(CH ₃) ₃ C |
| 813 | 4-F, 3-CH ₃ |
| 814 | 3-Cl, 4-Cl |
| 815 | 3,4-(CH ₂) ₄ |
| 816 | 3-HCF ₂ CF ₂ O |
| 817 | 3-CHF ₂ O |
| 818 | 3-(CH ₃) ₂ N |
| 819 | 3-cyclopropyl |
| 820 | 3-(2-furyl) |
| 821 | 3-CF ₃ CF ₂ |
| 822 | 4-NH ₂ |
| 823 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 824 | 4-CH ₃ CH ₂ CH ₂ O |

| Ex. No. | R _{SUB2} |
|---------|--|
| 827 | 3-CF ₃ O-benzyloxy |
| 828 | 3-CF ₃ -benzyloxy |
| 829 | 3-F, 5-F-benzyloxy |
| 830 | cyclohexylmethylenoxy |
| 831 | benzyloxy |
| 832 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 833 | 4-CF ₃ O-benzyloxy |
| 834 | 4-CH ₃ CH ₂ -benzyloxy |
| 835 | isopropoxy |
| 836 | 3-CF ₃ -benzyl |
| 837 | isopropylthio |
| 838 | cyclopentoxy |
| 839 | 3-Cl-5-pyridinyloxy |
| 840 | 3-CF ₃ S-benzyloxy |
| 841 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 842 | 2-F, 3-CF ₃ -benzyloxy |
| 843 | 3-F, 5-CF ₃ -benzyloxy |
| 844 | 4-(CH ₃) ₂ CH-benzyloxy |
| 845 | 1-phenylethoxy |
| 846 | 4-F, 3-CH ₃ -benzoyl |
| 847 | 3-CF ₃ -phenyl |
| 848 | 4-CH ₃ O-phenylamino |

Example Table 9. (3*R*)-4-[*N*-(aryl)-[(aryl)methyl]amino]-1,1,1,2,2-pentafluoro-3-butanols (Continued).

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|-------------------------|
| 825 | 3-CF ₃ |
| 826 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|-------------------------------|
| 849 | cyclopropoxy |
| 850 | 4-NO ₂ -phenylthio |



| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 851 | 3-isopropyl |
| 852 | 2-Cl, 3-Cl |
| 853 | 3-CF ₃ O |
| 854 | 4-F |
| 855 | 4-CH ₃ |
| 856 | 2-F, 5-Br |
| 857 | 4-Cl, 3-CH ₃ CH ₂ |
| 858 | 3-CH ₃ CH ₂ |
| 859 | 3-CH ₃ , 5-CH ₃ |
| 860 | 3-(CH ₃) ₃ C |
| 861 | 4-F, 3-CH ₃ |
| 862 | 3-Cl, 4-Cl |
| 863 | 3,4-(CH ₂) ₄ |
| 864 | 3-HCF ₂ CF ₂ O |
| 865 | 3-CHF ₂ O |
| 866 | 3-(CH ₃) ₂ N |
| 867 | 3-cyclopropyl |
| 868 | 3-(2-furyl) |

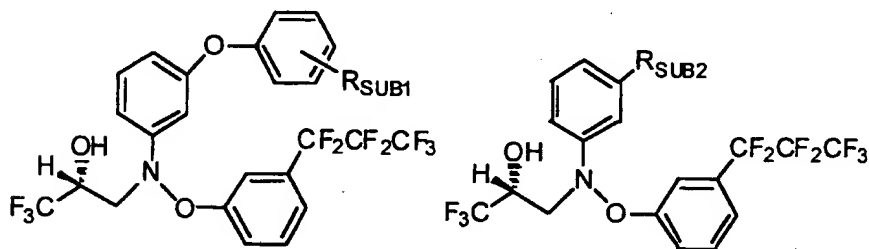
| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 875 | 3-CF ₃ O-benzyloxy |
| 876 | 3-CF ₃ -benzyloxy |
| 877 | 3-F, 5-F-benzyloxy |
| 878 | cyclohexylmethylenoxy |
| 879 | benzyloxy |
| 880 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 881 | 4-CF ₃ O-benzyloxy |
| 882 | 4-CH ₃ CH ₂ -benzyloxy |
| 883 | isopropoxy |
| 884 | 3-CF ₃ -benzyl |
| 885 | isopropylthio |
| 886 | cyclopentoxy |
| 887 | 3-Cl-5-pyridinyloxy |
| 888 | 3-CF ₃ S-benzyloxy |
| 889 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 890 | 2-F, 3-CF ₃ -benzyloxy |
| 891 | 3-F, 5-CF ₃ -benzyloxy |
| 892 | 4-(CH ₃) ₂ CH-benzyloxy |

Example Table 9. (3*R*)-4-[*N*-(aryl)-[(aryl)methylamino]-1,1,1,2,2-pentafluoro-3-butanols (Continued).

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 869 | 3-CF ₃ CF ₂ |
| 870 | 4-NH ₂ |
| 871 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 872 | 4-CH ₃ CH ₂ CH ₂ O |
| 873 | 3-CF ₃ |
| 874 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|---------------------------------|
| 893 | 1-phenylethoxy |
| 894 | 4-F, 3-CH ₃ -benzoyl |
| 895 | 3-CF ₃ -phenyl |
| 896 | 4-CH ₃ O-phenylamino |
| 897 | cyclopropoxy |
| 898 | 4-NO ₂ -phenylthio |

Example Table 10. Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)oxy]amino]-1,1,1-trifluoro-2-propanols.



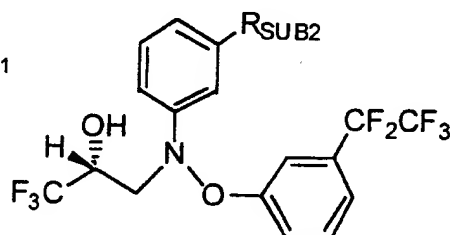
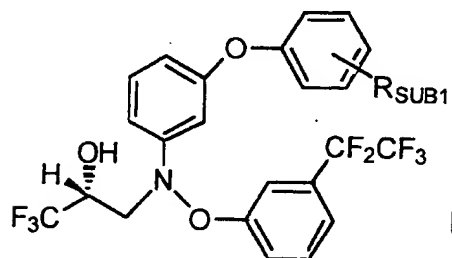
| Ex. No. | R _{SUB1} |
|---------|---|
| 899 | 3-isopropyl |
| 900 | 2-Cl, 3-Cl |
| 901 | 3-CF ₃ O |
| 902 | 4-F |
| 903 | 4-CH ₃ |
| 904 | 2-F, 5-Br |
| 905 | 4-Cl, 3-CH ₃ CH ₂ |
| 906 | 3-CH ₃ CH ₂ |
| 907 | 3-CH ₃ , 5-CH ₃ |
| 908 | 3-(CH ₃) ₃ C |
| 909 | 4-F, 3-CH ₃ |
| 910 | 3-Cl, 4-Cl |
| 911 | 3,4-(CH ₂) ₄ |
| 912 | 3-HCF ₂ CF ₂ O |
| 913 | 3-CHF ₂ O |
| 914 | 3-(CH ₃) ₂ N |
| 915 | 3-cyclopropyl |
| 916 | 3-(2-furyl) |
| 917 | 3-CF ₃ CF ₂ |
| 918 | 4-NH ₂ |
| 919 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 920 | 4-CH ₃ CH ₂ CH ₂ O |
| 921 | 3-CF ₃ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 923 | 3-CF ₃ O-benzyloxy |
| 924 | 3-CF ₃ -benzyloxy |
| 925 | 3-F, 5-F-benzyloxy |
| 926 | cyclohexylmethylenoxy |
| 927 | benzyloxy |
| 928 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 929 | 4-CF ₃ O-benzyloxy |
| 930 | 4-CH ₃ CH ₂ -benzyloxy |
| 931 | isopropoxy |
| 932 | 3-CF ₃ -benzyl |
| 933 | isopropylthio |
| 934 | cyclopentoxy |
| 935 | 3-Cl-5-pyridinyloxy |
| 936 | 3-CF ₃ S-benzyloxy |
| 937 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 938 | 2-F, 3-CF ₃ -benzyloxy |
| 939 | 3-F, 5-CF ₃ -benzyloxy |
| 940 | 4-(CH ₃) ₂ CH-benzyloxy |
| 941 | 1-phenylethoxy |
| 942 | 4-F, 3-CH ₃ -benzoyl |
| 943 | 3-CF ₃ -phenyl |
| 944 | 4-CH ₃ O-phenylamino |
| 945 | cyclopropoxy |

Example Table 10 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)oxy]amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|-------------------------|
| 922 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|-------------------------------|
| 946 | 4-NO ₂ -phenylthio |



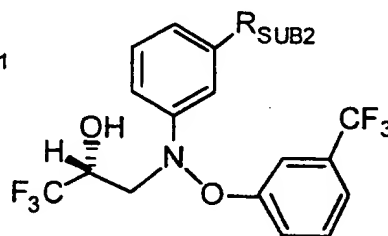
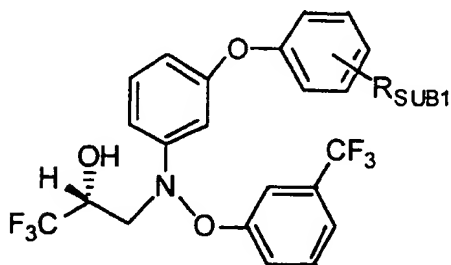
| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 947 | 3-isopropyl |
| 948 | 2-Cl, 3-Cl |
| 949 | 3-CF ₃ O |
| 950 | 4-F |
| 951 | 4-CH ₃ |
| 952 | 2-F, 5-Br |
| 953 | 4-Cl, 3-CH ₃ CH ₂ |
| 954 | 3-CH ₃ CH ₂ |
| 955 | 3-CH ₃ , 5-CH ₃ |
| 956 | 3-(CH ₃) ₃ C |
| 957 | 4-F, 3-CH ₃ |
| 958 | 3-Cl, 4-Cl |
| 959 | 3,4-(CH ₂) ₄ |
| 960 | 3-HCF ₂ CF ₂ O |
| 961 | 3-CHF ₂ O |
| 962 | 3-(CH ₃) ₂ N |
| 963 | 3-cyclopropyl |
| 964 | 3-(2-furyl) |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 971 | 3-CF ₃ O-benzyloxy |
| 972 | 3-CF ₃ -benzyloxy |
| 973 | 3-F, 5-F-benzyloxy |
| 974 | cyclohexylmethylenoxy |
| 975 | benzyloxy |
| 976 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 977 | 4-CF ₃ O-benzyloxy |
| 978 | 4-CH ₃ CH ₂ -benzyloxy |
| 979 | isopropoxy |
| 980 | 3-CF ₃ -benzyl |
| 981 | isopropylthio |
| 982 | cyclopentoxy |
| 983 | 3-Cl-5-pyridinyloxy |
| 984 | 3-CF ₃ S-benzyloxy |
| 985 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 986 | 2-F, 3-CF ₃ -benzyloxy |
| 987 | 3-F, 5-CF ₃ -benzyloxy |
| 988 | 4-(CH ₃) ₂ CH-benzyloxy |

Example Table 10 (continued). Substituted (2*R*)-3-[*N*-(aryl)-|(aryl)oxy|amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 965 | 3-CF ₃ CF ₂ |
| 966 | 4-NH ₂ |
| 967 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 968 | 4-CH ₃ CH ₂ CH ₂ O |
| 969 | 3-CF ₃ |
| 970 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|---------------------------------|
| 989 | 1-phenylethoxy |
| 990 | 4-F, 3-CH ₃ -benzoyl |
| 991 | 3-CF ₃ -phenyl |
| 992 | 4-CH ₃ O-phenylamino |
| 993 | cyclopropoxy |
| 994 | 4-NO ₂ -phenylthio |



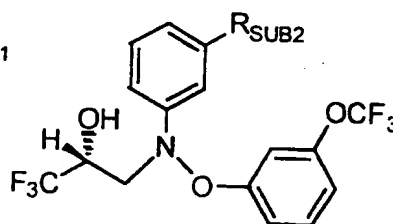
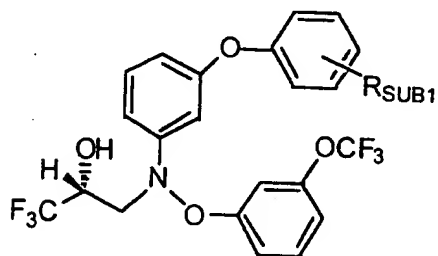
| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 995 | 3-isopropyl |
| 996 | 2-Cl, 3-Cl |
| 997 | 3-CF ₃ O |
| 998 | 4-F |
| 999 | 4-CH ₃ |
| 1000 | 2-F, 5-Br |
| 1001 | 4-Cl, 3-CH ₃ CH ₂ |
| 1002 | 3-CH ₃ CH ₂ |
| 1003 | 3-CH ₃ , 5-CH ₃ |
| 1004 | 3-(CH ₃) ₃ C |
| 1005 | 4-F, 3-CH ₃ |
| 1006 | 3-Cl, 4-Cl |
| 1007 | 3,4-(CH ₂) ₄ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 1019 | 3-CF ₃ O-benzyloxy |
| 1020 | 3-CF ₃ -benzyloxy |
| 1021 | 3-F, 5-F-benzyloxy |
| 1022 | cyclohexylmethylenoxy |
| 1023 | benzyloxy |
| 1024 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1025 | 4-CF ₃ O-benzyloxy |
| 1026 | 4-CH ₃ CH ₂ -benzyloxy |
| 1027 | isopropoxy |
| 1028 | 3-CF ₃ -benzyl |
| 1029 | isopropylthio |
| 1030 | cyclopentoxy |
| 1031 | 3-Cl-5-pyridinyloxy |

Example Table 10 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)oxy]amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 1008 | 3-HCF ₂ CF ₂ O |
| 1009 | 3-CHF ₂ O |
| 1010 | 3-(CH ₃) ₂ N |
| 1011 | 3-cyclopropyl |
| 1012 | 3-(2-furyl) |
| 1013 | 3-CF ₃ CF ₂ |
| 1014 | 4-NH ₂ |
| 1015 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1016 | 4-CH ₃ CH ₂ CH ₂ O |
| 1017 | 3-CF ₃ |
| 1018 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 1032 | 3-CF ₃ S-benzyloxy |
| 1033 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1034 | 2-F, 3-CF ₃ -benzyloxy |
| 1035 | 3-F, 5-CF ₃ -benzyloxy |
| 1036 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1037 | 1-phenylethoxy |
| 1038 | 4-F, 3-CH ₃ -benzoyl |
| 1039 | 3-CF ₃ -phenyl |
| 1040 | 4-CH ₃ O-phenylamino |
| 1041 | cyclopropoxy |
| 1042 | 4-NO ₂ -phenylthio |



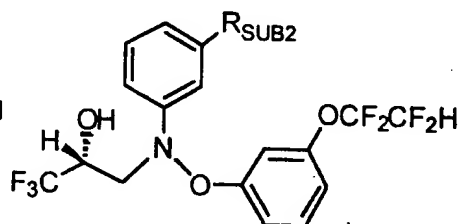
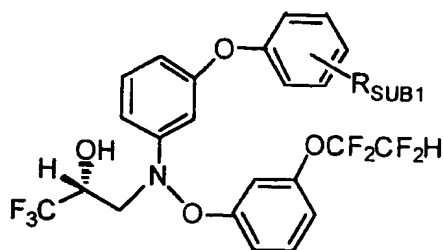
| Ex. No. | R _{SUB1} |
|---------|---|
| 1043 | 3-isopropyl |
| 1044 | 2-Cl, 3-Cl |
| 1045 | 3-CF ₃ O |
| 1046 | 4-F |
| 1047 | 4-CH ₃ |
| 1048 | 2-F, 5-Br |
| 1049 | 4-Cl, 3-CH ₃ CH ₂ |
| 1050 | 3-CH ₃ CH ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 1067 | 3-CF ₃ O-benzyloxy |
| 1068 | 3-CF ₃ -benzyloxy |
| 1069 | 3-F, 5-F-benzyloxy |
| 1070 | cyclohexylmethylenoxy |
| 1071 | benzyloxy |
| 1072 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1073 | 4-CF ₃ O-benzyloxy |
| 1074 | 4-CH ₃ CH ₂ -benzyloxy |

Example Table 10 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)oxy|amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 1051 | 3-CH ₃ , 5-CH ₃ |
| 1052 | 3-(CH ₃) ₃ C |
| 1053 | 4-F, 3-CH ₃ |
| 1054 | 3-Cl, 4-Cl |
| 1055 | 3,4-(CH ₂) ₄ |
| 1056 | 3-HCF ₂ CF ₂ O |
| 1057 | 3-CHF ₂ O |
| 1058 | 3-(CH ₃) ₂ N |
| 1059 | 3-cyclopropyl |
| 1060 | 3-(2-furyl) |
| 1061 | 3-CF ₃ CF ₂ |
| 1062 | 4-NH ₂ |
| 1063 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1064 | 4-CH ₃ CH ₂ CH ₂ O |
| 1065 | 3-CF ₃ |
| 1066 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 1075 | isopropoxy |
| 1076 | 3-CF ₃ -benzyl |
| 1077 | isopropylthio |
| 1078 | cyclopentoxy |
| 1079 | 3-Cl-5-pyridinyloxy |
| 1080 | 3-CF ₃ S-benzyloxy |
| 1081 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1082 | 2-F, 3-CF ₃ -benzyloxy |
| 1083 | 3-F, 5-CF ₃ -benzyloxy |
| 1084 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1085 | 1-phenylethoxy |
| 1086 | 4-F, 3-CH ₃ -benzoyl |
| 1087 | 3-CF ₃ -phenyl |
| 1088 | 4-CH ₃ O-phenylamino |
| 1089 | cyclopropoxy |
| 1090 | 4-NO ₂ -phenylthio |



| Ex. No. | R _{SUB1} |
|---------|---------------------|
| 1091 | 3-isopropyl |
| 1092 | 2-Cl, 3-Cl |
| 1093 | 3-CF ₃ O |
| 1094 | 4-F |

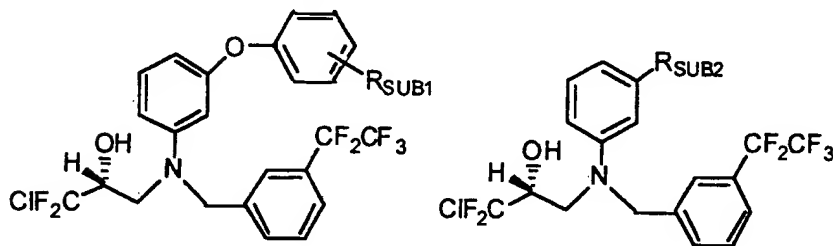
| Ex. No. | R _{SUB2} |
|---------|-------------------------------|
| 1115 | 3-CF ₃ O-benzyloxy |
| 1116 | 3-CF ₃ -benzyloxy |
| 1117 | 3-F, 5-F-benzyloxy |
| 1118 | cyclohexylmethylenoxy |

Example Table 10 (continued). Substituted (2*R*)-3-[*N*-(aryl)-[(aryl)oxy]amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 1095 | 4-CH ₃ |
| 1096 | 2-F, 5-Br |
| 1097 | 4-Cl, 3-CH ₃ CH ₂ |
| 1098 | 3-CH ₃ CH ₂ |
| 1099 | 3-CH ₃ , 5-CH ₃ |
| 1100 | 3-(CH ₃) ₃ C |
| 1101 | 4-F, 3-CH ₃ |
| 1102 | 3-Cl, 4-Cl |
| 1103 | 3,4-(CH ₂) ₄ |
| 1104 | 3-HCF ₂ CF ₂ O |
| 1105 | 3-CHF ₂ O |
| 1106 | 3-(CH ₃) ₂ N |
| 1107 | 3-cyclopropyl |
| 1108 | 3-(2-furyl) |
| 1109 | 3-CF ₃ CF ₂ |
| 1110 | 4-NH ₂ |
| 1111 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1112 | 4-CH ₃ CH ₂ CH ₂ O |
| 1113 | 3-CF ₃ |
| 1114 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|--|
| 1119 | benzyloxy |
| 1120 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1121 | 4-CF ₃ O-benzyloxy |
| 1122 | 4-CH ₃ CH ₂ -benzyloxy |
| 1123 | isopropoxy |
| 1124 | 3-CF ₃ -benzyl |
| 1125 | isopropylthio |
| 1126 | cyclopentoxy |
| 1127 | 3-Cl-5-pyridinyloxy |
| 1128 | 3-CF ₃ S-benzyloxy |
| 1129 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1130 | 2-F, 3-CF ₃ -benzyloxy |
| 1131 | 3-F, 5-CF ₃ -benzyloxy |
| 1132 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1133 | 1-phenylethoxy |
| 1134 | 4-F, 3-CH ₃ -benzoyl |
| 1135 | 3-CF ₃ -phenyl |
| 1136 | 4-CH ₃ O-phenylamino |
| 1137 | cyclopropoxy |
| 1138 | 4-NO ₂ -phenylthio |

Example Table 11. (2*R*)-3-[*N*-(aryl)-[(aryl)methyl]amino]-1,1-difluoro-1-chloro-2-propanols.



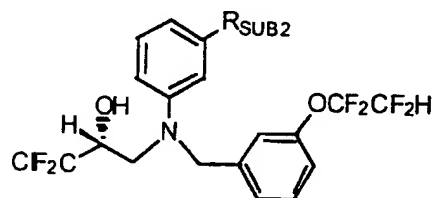
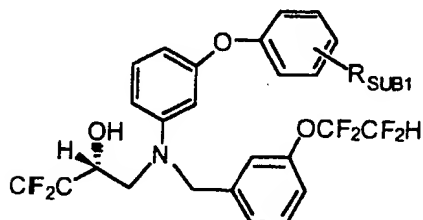
| Ex. No. | R _{SUB1} |
|---------|---|
| 1139 | 3-isopropyl |
| 1140 | 2-Cl, 3-Cl |
| 1141 | 3-CF ₃ O |
| 1142 | 4-F |
| 1143 | 4-CH ₃ |
| 1144 | 2-F, 5-Br |
| 1145 | 4-Cl, 3-CH ₃ CH ₂ |
| 1146 | 3-CH ₃ CH ₂ |
| 1147 | 3-CH ₃ , 5-CH ₃ |
| 1148 | 3-(CH ₃) ₃ C |
| 1149 | 4-F, 3-CH ₃ |
| 1150 | 3-Cl, 4-Cl |
| 1151 | 3,4-(CH ₂) ₄ |
| 1152 | 3-HCF ₂ CF ₂ O |
| 1153 | 3-CHF ₂ O |
| 1154 | 3-(CH ₃) ₂ N |
| 1155 | 3-cyclopropyl |
| 1156 | 3-(2-furyl) |
| 1157 | 3-CF ₃ CF ₂ |
| 1158 | 4-NH ₂ |
| 1159 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1160 | 4-CH ₃ CH ₂ CH ₂ O |
| 1161 | 3-CF ₃ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 1163 | 3-CF ₃ O-benzyloxy |
| 1164 | 3-CF ₃ -benzyloxy |
| 1165 | 3-F, 5-F-benzyloxy |
| 1166 | cyclohexylmethylenoxy |
| 1167 | benzyloxy |
| 1168 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1169 | 4-CF ₃ O-benzyloxy |
| 1170 | 4-CH ₃ CH ₂ -benzyloxy |
| 1171 | isopropoxy |
| 1172 | 3-CF ₃ -benzyl |
| 1173 | isopropylthio |
| 1174 | cyclopentoxy |
| 1175 | 3-Cl-5-pyridinyloxy |
| 1176 | 3-CF ₃ S-benzyloxy |
| 1177 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1178 | 2-F, 3-CF ₃ -benzyloxy |
| 1179 | 3-F, 5-CF ₃ -benzyloxy |
| 1180 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1181 | 1-phenylethoxy |
| 1182 | 4-F, 3-CH ₃ -benzoyl |
| 1183 | 3-CF ₃ -phenyl |
| 1184 | 4-CH ₃ O-phenylamino |
| 1185 | cyclopropoxy |

Example Table 11 (continued). (2*R*)-3-[*N*-(aryl)-[(aryl)methyl]amino]-1,1-difluoro-1-chloro-2-propanols.

| Ex. No. | R _{SUB1} |
|------------|-------------------|
| 1162 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|------------|-------------------------------|
| 1186 | 4-NO ₂ -phenylthio |



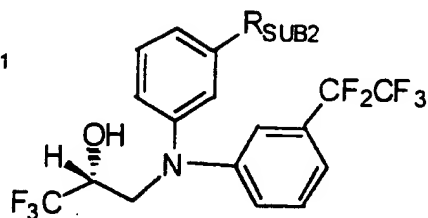
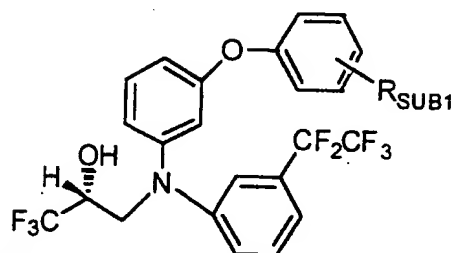
| Ex. No. | R _{SUB1} |
|------------|---|
| 1187 | 3-isopropyl |
| 1188 | 2-Cl, 3-Cl |
| 1189 | 3-CF ₃ O |
| 1190 | 4-F |
| 1191 | 4-CH ₃ |
| 1192 | 2-F, 5-Br |
| 1193 | 4-Cl, 3-CH ₃ CH ₂ |
| 1194 | 3-CH ₃ CH ₂ |
| 1195 | 3-CH ₃ , 5-CH ₃ |
| 1196 | 3-(CH ₃) ₃ C |
| 1197 | 4-F, 3-CH ₃ |
| 1198 | 3-Cl, 4-Cl |
| 1199 | 3,4-(CH ₂) ₄ |
| 1200 | 3-HCF ₂ CF ₂ O |
| 1201 | 3-CHF ₂ O |
| 1202 | 3-(CH ₃) ₂ N |
| 1203 | 3-cyclopropyl |
| 1204 | 3-(2-furyl) |
| 1205 | 3-CF ₃ CF ₂ |

| Ex. No. | R _{SUB2} |
|------------|--|
| 1211 | 3-CF ₃ O-benzyloxy |
| 1212 | 3-CF ₃ -benzyloxy |
| 1213 | 3-F, 5-F-benzyloxy |
| 1214 | cyclohexylmethylenedioxy |
| 1215 | benzyloxy |
| 1216 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1217 | 4-CF ₃ O-benzyloxy |
| 1218 | 4-CH ₃ CH ₂ -benzyloxy |
| 1219 | isopropoxy |
| 1220 | 3-CF ₃ -benzyl |
| 1221 | isopropylthio |
| 1222 | cyclopentoxy |
| 1223 | 3-Cl-5-pyridinyloxy |
| 1224 | 3-CF ₃ S-benzyloxy |
| 1225 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1226 | 2-F, 3-CF ₃ -benzyloxy |
| 1227 | 3-F, 5-CF ₃ -benzyloxy |
| 1228 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1229 | 1-phenylethoxy |

Example Table 11 (continued). (2*R*)-3-[*N*-(aryl)-[(aryl)methyl]amino]-1,1-difluoro-1-chloro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 1206 | 4-NH ₂ |
| 1207 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1208 | 4-CH ₃ CH ₂ CH ₂ O |
| 1209 | 3-CF ₃ |
| 1210 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|---------------------------------|
| 1230 | 4-F, 3-CH ₃ -benzoyl |
| 1231 | 3-CF ₃ -phenyl |
| 1232 | 4-CH ₃ O-phenylamino |
| 1233 | cyclopropoxy |
| 1234 | 4-NO ₂ -phenylthio |

Example Table 12. (2*R*)-3-[*N,N'*-(diaryl)amino]-1,1,1-trifluoro-2-propanols.

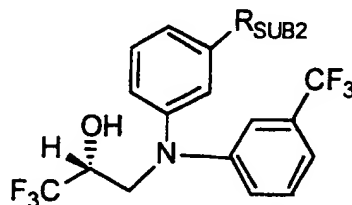
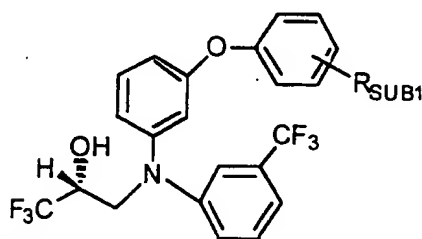
| Ex. No. | R_{SUB1} |
|---------|--|
| 1235 | 3-isopropyl |
| 1236 | 2-Cl, 3-Cl |
| 1237 | 3- CF_3O |
| 1238 | 4-F |
| 1239 | 4- CH_3 |
| 1240 | 2-F, 5-Br |
| 1241 | 4-Cl, 3- CH_3CH_2 |
| 1242 | 3- CH_3CH_2 |
| 1243 | 3- CH_3 , 5- CH_3 |
| 1244 | 3-(CH_3) $_3\text{C}$ |
| 1245 | 4-F, 3- CH_3 |
| 1246 | 3-Cl, 4-Cl |
| 1247 | 3,4-(CH_2) $_4$ |
| 1248 | 3- $\text{HCF}_2\text{CF}_2\text{O}$ |
| 1249 | 3- CHF_2O |
| 1250 | 3-(CH_3) $_2\text{N}$ |
| 1251 | 3-cyclopropyl |
| 1252 | 3-(2-furyl) |
| 1253 | 3- CF_3CF_2 |
| 1254 | 4- NH_2 |
| 1255 | 3- CH_3 , 4- CH_3 , 5- CH_3 |
| 1256 | 4- $\text{CH}_3\text{CH}_2\text{CH}_2\text{O}$ |
| 1257 | 3- CF_3 |

| Ex. No. | R_{SUB2} |
|---------|--|
| 1259 | 3- CF_3O -benzyloxy |
| 1260 | 3- CF_3 -benzyloxy |
| 1261 | 3-F, 5-F-benzyloxy |
| 1262 | cyclohexylmethylenedioxy |
| 1263 | benzyloxy |
| 1264 | 3- CF_3 , 5- CF_3 -benzyloxy |
| 1265 | 4- CF_3O -benzyloxy |
| 1266 | 4- CH_3CH_2 -benzyloxy |
| 1267 | isopropoxy |
| 1268 | 3- CF_3 -benzyl |
| 1269 | isopropylthio |
| 1270 | cyclopentoxy |
| 1271 | 3-Cl-5-pyridinyloxy |
| 1272 | 3- CF_3S -benzyloxy |
| 1273 | 3- CH_3 , 4- CH_3 -benzyloxy |
| 1274 | 2-F, 3- CF_3 -benzyloxy |
| 1275 | 3-F, 5- CF_3 -benzyloxy |
| 1276 | 4-(CH_3) $_2\text{CH}$ -benzyloxy |
| 1277 | 1-phenylethoxy |
| 1278 | 4-F, 3- CH_3 -benzoyl |
| 1279 | 3- CF_3 -phenyl |
| 1280 | 4- CH_3O -phenylamino |
| 1281 | cyclopropoxy |

Example Table 12 (continued). (2*R*)-3-[*N,N'*-(diaryl)amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|-------------------------|
| 1258 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|-------------------------------|
| 1282 | 4-NO ₂ -phenylthio |



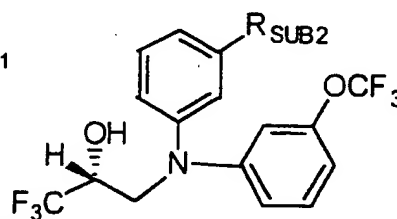
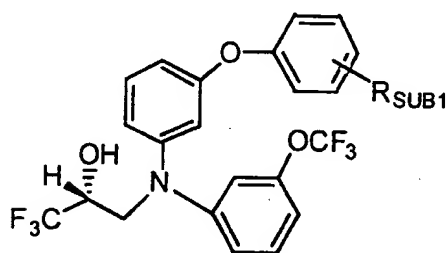
| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|----------------|---|
| 1283 | 3-isopropyl |
| 1284 | 2-Cl, 3-Cl |
| 1285 | 3-CF ₃ O |
| 1286 | 4-F |
| 1287 | 4-CH ₃ |
| 1288 | 2-F, 5-Br |
| 1289 | 4-Cl, 3-CH ₃ CH ₂ |
| 1290 | 3-CH ₃ CH ₂ |
| 1291 | 3-CH ₃ , 5-CH ₃ |
| 1292 | 3-(CH ₃) ₃ C |
| 1293 | 4-F, 3-CH ₃ |
| 1294 | 3-Cl, 4-Cl |
| 1295 | 3,4-(CH ₂) ₄ |
| 1296 | 3-HCF ₂ CF ₂ O |
| 1297 | 3-CHF ₂ O |
| 1298 | 3-(CH ₃) ₂ N |
| 1299 | 3-cyclopropyl |
| 1300 | 3-(2-furyl) |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|----------------|--|
| 1307 | 3-CF ₃ O-benzyloxy |
| 1308 | 3-CF ₃ -benzyloxy |
| 1309 | 3-F, 5-F-benzyloxy |
| 1310 | cyclohexylmethyleneoxy |
| 1311 | benzyloxy |
| 1312 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1313 | 4-CF ₃ O-benzyloxy |
| 1314 | 4-CH ₃ CH ₂ -benzyloxy |
| 1315 | isopropoxy |
| 1316 | 3-CF ₃ -benzyl |
| 1317 | isopropylthio |
| 1318 | cyclopentoxy |
| 1319 | 3-Cl-5-pyridinyloxy |
| 1320 | 3-CF ₃ S-benzyloxy |
| 1321 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1322 | 2-F, 3-CF ₃ -benzyloxy |
| 1323 | 3-F, 5-CF ₃ -benzyloxy |
| 1324 | 4-(CH ₃) ₂ CH-benzyloxy |

Example Table 12 (continued). (2*R*)-3-[*N,N'*-(diaryl)amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 1301 | 3-CF ₃ CF ₂ |
| 1302 | 4-NH ₂ |
| 1303 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1304 | 4-CH ₃ CH ₂ CH ₂ O |
| 1305 | 3-CF ₃ |
| 1306 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|---------------------------------|
| 1325 | 1-phenylethoxy |
| 1326 | 4-F, 3-CH ₃ -benzoyl |
| 1327 | 3-CF ₃ -phenyl |
| 1328 | 4-CH ₃ O-phenylamino |
| 1329 | cyclopropoxy |
| 1330 | 4-NO ₂ -phenylthio |



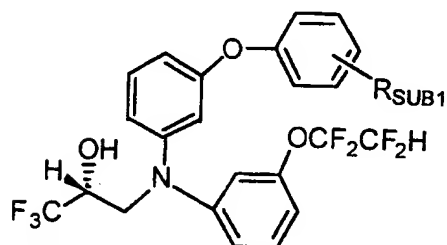
| Ex. No. | R _{SUB1} |
|---------|---|
| 1331 | 3-isopropyl |
| 1332 | 2-Cl, 3-Cl |
| 1333 | 3-CF ₃ O |
| 1334 | 4-F |
| 1335 | 4-CH ₃ |
| 1336 | 2-F, 5-Br |
| 1337 | 4-Cl, 3-CH ₃ CH ₂ |
| 1338 | 3-CH ₃ CH ₂ |
| 1339 | 3-CH ₃ , 5-CH ₃ |
| 1340 | 3-(CH ₃) ₃ C |
| 1341 | 4-F, 3-CH ₃ |
| 1342 | 3-Cl, 4-Cl |
| 1343 | 3,4-(CH ₂) ₄ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 1355 | 3-CF ₃ O-benzyloxy |
| 1356 | 3-CF ₃ -benzyloxy |
| 1357 | 3-F, 5-F-benzyloxy |
| 1358 | cyclohexylmethylenedioxy |
| 1359 | benzyloxy |
| 1360 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1361 | 4-CF ₃ O-benzyloxy |
| 1362 | 4-CH ₃ CH ₂ -benzyloxy |
| 1363 | isopropoxy |
| 1364 | 3-CF ₃ -benzyl |
| 1365 | isopropylthio |
| 1366 | cyclopentoxy |
| 1367 | 3-Cl-5-pyridinyloxy |

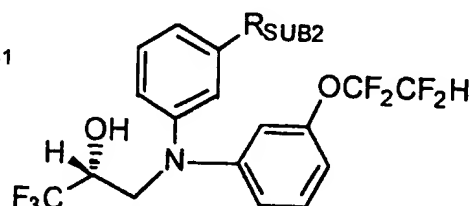
Example Table 12 (continued). (2*R*)-3-[*N,N'*-(diaryl)amino]-1,1,1-trifluoro-2-propanols.

| Ex. No. | R _{SUB1} |
|---------|---|
| 1344 | 3-HCF ₂ CF ₂ O |
| 1345 | 3-CHF ₂ O |
| 1346 | 3-(CH ₃) ₂ N |
| 1347 | 3-cyclopropyl |
| 1348 | 3-(2-furyl) |
| 1349 | 3-CF ₃ CF ₂ |
| 1350 | 4-NH ₂ |
| 1351 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1352 | 4-CH ₃ CH ₂ CH ₂ O |
| 1353 | 3-CF ₃ |
| 1354 | 2-NO ₂ |

| Ex. No. | R _{SUB2} |
|---------|--|
| 1368 | 3-CF ₃ S-benzyloxy |
| 1369 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1370 | 2-F, 3-CF ₃ -benzyloxy |
| 1371 | 3-F, 5-CF ₃ -benzyloxy |
| 1372 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1373 | 1-phenylethoxy |
| 1374 | 4-F, 3-CH ₃ -benzoyl |
| 1375 | 3-CF ₃ -phenyl |
| 1376 | 4-CH ₃ O-phenylamino |
| 1377 | cyclopropoxy |
| 1378 | 4-NO ₂ -phenylthio |



| Ex. No. | R _{SUB1} |
|---------|---|
| 1379 | 3-isopropyl |
| 1380 | 2-Cl, 3-Cl |
| 1381 | 3-CF ₃ O |
| 1382 | 4-F |
| 1383 | 4-CH ₃ |
| 1384 | 2-F, 5-Br |
| 1385 | 4-Cl, 3-CH ₃ CH ₂ |
| 1386 | 3-CH ₃ CH ₂ |



| Ex. No. | R _{SUB2} |
|---------|--|
| 1403 | 3-CF ₃ O-benzyloxy |
| 1404 | 3-CF ₃ -benzyloxy |
| 1405 | 3-F, 5-F-benzyloxy |
| 1406 | cyclohexylmethylenoxy |
| 1407 | benzyloxy |
| 1408 | 3-CF ₃ , 5-CF ₃ -benzyloxy |
| 1409 | 4-CF ₃ O-benzyloxy |
| 1410 | 4-CH ₃ CH ₂ -benzyloxy |

Example Table 12 (continued). (2*R*)-3-[*N,N'*-(diaryl)amino]-1,1,1-trifluoro-2-propanols.

| <u>Ex. No.</u> | <u>R_{SUB1}</u> |
|--------------------|---|
| 1387 | 3-CH ₃ , 5-CH ₃ |
| 1388 | 3-(CH ₃) ₃ C |
| 1389 | 4-F, 3-CH ₃ |
| 1390 | 3-Cl, 4-Cl |
| 1391 | 3,4-(CH ₂) ₄ |
| 1392 | 3-HCF ₂ CF ₂ O |
| 1393 | 3-CHF ₂ O |
| 1394 | 3-(CH ₃) ₂ N |
| 1395 | 3-cyclopropyl |
| 1396 | 3-(2-furyl) |
| 1397 | 3-CF ₃ CF ₂ |
| 1398 | 4-NH ₂ |
| 1399 | 3-CH ₃ , 4-CH ₃ , 5-CH ₃ |
| 1400 | 4-CH ₃ CH ₂ CH ₂ O |
| 1401 | 3-CF ₃ |
| 1402 | 2-NO ₂ |

| <u>Ex. No.</u> | <u>R_{SUB2}</u> |
|--------------------|--|
| 1411 | isopropoxy |
| 1412 | 3-CF ₃ -benzyl |
| 1413 | isopropylthio |
| 1414 | cyclopentoxy |
| 1415 | 3-Cl-5-pyridinyloxy |
| 1416 | 3-CF ₃ S-benzyloxy |
| 1417 | 3-CH ₃ , 4-CH ₃ -benzyloxy |
| 1418 | 2-F, 3-CF ₃ -benzyloxy |
| 1419 | 3-F, 5-CF ₃ -benzyloxy |
| 1420 | 4-(CH ₃) ₂ CH-benzyloxy |
| 1421 | 1-phenylethoxy |
| 1422 | 4-F, 3-CH ₃ -benzoyl |
| 1423 | 3-CF ₃ -phenyl |
| 1424 | 4-CH ₃ O-phenylamino |
| 1425 | cyclopropoxy |
| 1426 | 4-NO ₂ -phenylthio |

BIOASSAYS

CETP Activity *In Vitro*ASSAY OF CETP INHIBITION USING PURIFIED COMPONENTS
(RECONSTITUTED BUFFER ASSAY)

- 5 The ability of compounds to inhibit CETP activity was assessed using an *in vitro* assay that measured the rate of transfer of radiolabeled cholesteryl ester ([³H]CE) from HDL donor particles to LDL acceptor particles. Details of the assay are provided by Glenn, K. C. et al. (Glenn and Melton, "Quantification of Cholesteryl Ester Transfer Protein (CETP): A) CETP Activity and B)
- 10 Immunochemical Assay of CETP Protein," *Meth. Enzymol.*, 263, 339-351 (1996)). Human recombinant CETP can be obtained from the serum-free conditioned medium of CHO cells transfected with a cDNA for CETP and purified as described by Wang, S. et al. (*J. Biol. Chem.* 267, 17487-17490 (1992)). To measure CETP activity, [³H]CE-labeled-HDL, LDL, CETP and
- 15 assay buffer (50 mM tris(hydroxymethyl)aminomethane, pH 7.4; 150 mM sodium chloride; 2 mM ethylenediamine-tetraacetic acid (EDTA); 1% bovine serum albumin) were incubated in a final volume of 200 μ L, for 2 hours at 37 °C in 96 well plates. Inhibitors were included in the assay by diluting from a 10 mM DMSO stock solution into 16% (v/v) aqueous DMSO so that the final
- 20 concentration of inhibitor was 800 μ M. The inhibitors were then diluted 1:1 with CETP in assay buffer, and then 25 μ L of that solution was mixed with 175 μ L of lipoprotein pool for assay. Following incubation, LDL was differentially precipitated by the addition of 50 μ L of 1% (w/v) dextran sulfate/0.5 M magnesium chloride, mixed by vortex, and incubated at room temperature for 10
- 25 minutes. A portion of the solution (200 μ L) was transferred to a filter plate (Millipore). After filtration, the radioactivity present in the precipitated LDL was measured by liquid scintillation counting. Correction for non-specific transfer or precipitation was made by including samples that do not contain CETP. The rate

of [^3H]CE transfer using this assay was linear with respect to time and CETP concentration. up to 25-30% of [^3H]CE transferred.

The potency of test compounds was determined by performing the above described assay in the presence of varying concentrations of the test compounds
5 and determining the concentration required for 50% inhibition of transfer of [^3H]CE from HDL to LDL. This value was defined as the IC_{50} . The IC_{50} values determined from this assay are accurate when the IC_{50} is greater than 10 nM. In the case where compounds have greater inhibitory potency, accurate measurements of IC_{50} may be determined using longer incubation times (up to
10 18 hours) and lower final concentrations of CETP (< 50 nM).

Examples of IC_{50} values determined by these methods are summarized in Table 9.

ASSAY OF CETP INHIBITION IN HUMAN PLASMA

Blood was obtained from healthy volunteers, recruited from the personnel
15 of Monsanto Company, Saint Louis, MO. Blood was collected in tubes containing EDTA (EDTA plasma pool). The EDTA human plasma pool, previously stored at -20 °C, was thawed at room temperature and centrifuged for 5 minutes to remove any particulate matter. Tritiated HDL, radiolabeled in the cholesteryl ester moiety ([^3H]CE-HDL) as described by Morton and Zilversmit
20 (J. Biol. Chem., 256, 11992-95 (1981)), was added to the plasma to a final concentration of 25 $\mu\text{g/mL}$ cholesterol. Equal volumes (396 μL) of the plasma containing the [^3H]CE-HDL were added by pipette into micro tubes (Titertube®, Bio-Rad laboratories, Hercules, CA). Inhibitor compounds, dissolved as 20-50 mM stock solutions in DMSO, were serially diluted in DMSO (or an alternative
25 solvent in some cases, such as dimethylformamide or ethanol). Four μL of each of the serial dilutions of inhibitor compounds or DMSO alone were then added to each of the tubes containing plasma (396 μL). After mixing, triplicate aliquots (100 μL) from each plasma tube were then transferred to wells of 96-well round-

bottomed polystyrene microtiter plates (Corning, Corning, NY). Plates were sealed with plastic film and incubated at 37 °C for 4 hours. "Test" samples contained plasma with dilutions of inhibitor compounds. "Control" samples contained plasma with DMSO diluted to the same concentration as the test samples, but without inhibitor. "Blank" samples were prepared as "control" samples, but were left in the micro tubes at 4 °C for the 4 hour incubation and were then added to the microtiter wells at the end of the incubation period. VLDL and LDL were precipitated by the addition of 10 µL of precipitating reagent (1% (w/v) dextran sulfate (Dextralip50)/0.5 M magnesium chloride, pH 7.4) to all wells. The wells were mixed on a plate mixer and then incubated at ambient temperature for 10 min. The plates were then centrifuged at 1000 x g for 30 min at 10 °C. The supernatants (50 µL) from each well were then transferred to Picoplate™ 96 plate wells (Packard, Meriden, CT) containing Microscint™-40 (Packard, Meriden, CT). The plates were heat-sealed (TopSeal™-P, Packard, Meriden, CT) according to the manufacturer's directions and mixed for 30 min. Radioactivity was measured on a microplate scintillation counter (TopCount, Packard, Meriden, CT). The maximum percentage transfer in the control wells (% transfer) was determined using the following equation:

$$\% \text{ Transfer} = \frac{[\text{dpm}_{\text{blank}} - \text{dpm}_{\text{control}}] \times 100}{\text{dpm}_{\text{blank}}}$$

The percentage of transfer relative to the control (% control) was determined in the wells containing inhibitor compounds was determined as follows:

$$\% \text{ Control} = \frac{[\text{dpm}_{\text{blank}} - \text{dpm}_{\text{test}}] \times 100}{\text{dpm}_{\text{blank}} - \text{dpm}_{\text{control}}}$$

IC₅₀ values were then calculated from plots of % control versus concentration of inhibitor compound. IC₅₀ values were determined as the concentration of inhibitor compound inhibiting transfer of [³H]CE from the supernatant [³H]CE-

HDL to the precipitated VLDL and LDL by 50% compared to the transfer obtained in the control wells.

Examples of plasma IC₅₀ values determined by these methods are summarized in Table 10.

5 ASSAY OF CETP INHIBITION *IN VIVO*.

Inhibition of CETP activity by a test compound can be determined by administering the compound to an animal by intravenous injection or oral gavage, measuring the amount of transfer of tritium-labeled cholesteryl ester ([³H]CE) from HDL to VLDL and LDL particles, and comparing this amount of transfer
10 with the amount of transfer observed in control animals.

Male golden Syrian hamsters were maintained on a diet of chow containing 0.24% cholesterol for at least two weeks prior to the study. For animals receiving intravenous dosing immediately before the experiment, animals were anesthetized with pentobarbital. Anesthesia was maintained throughout the
15 experiment. In-dwelling catheters were inserted into the jugular vein and carotid artery. At the start of the experiment all animals received 0.2 mL of a solution containing [³H]CE-HDL into the jugular vein. [³H]CE-HDL is a preparation of human HDL containing tritium-labeled cholesteryl ester, and was prepared according to the method of Glenn et al. (*Meth. Enzymol.*, 263, 339-351 (1996)).
20 Test compound was dissolved as a 80 mM stock solution in vehicle (2% ethanol: 98% PEG 400, Sigma Chemical Company, St. Louis, Missouri, USA) and administered either by bolus injection or by continuous infusion. Two minutes after the [³H]CE-HDL dose was administered, animals received 0.1 mL of the test solution injected into the jugular vein. Control animals received 0.1 mL of
25 the intravenous vehicle solution without test compound. After 5 minutes, the first blood samples (0.5 mL) were taken from the carotid artery and collected in standard microtainer tubes containing ethylenediamine tetraacetic acid. Saline (0.5 mL) was injected to flush the catheter and replace blood volume. Subsequent blood samples were taken at two hours and four hours by the same

method. Blood samples were mixed well and kept on ice until the completion of the experiment. Plasma was obtained by centrifugation of the blood samples at 4 °C. The plasma (50 µL) was treated with 5 µL of precipitating reagent (dextran sulfate, 10 g/L; 0.5 M magnesium chloride) to remove VLDL/LDL. After
5 centrifugation, the resulting supernatant (25 µL) containing the HDL was analyzed for radioactivity using a liquid scintillation counter.

The percentage [^3H]CE transferred from HDL to LDL and VLDL (% transfer) was calculated based on the total radioactivity in equivalent plasma samples before precipitation. Typically, the amount of transfer from HDL to
10 LDL and VLDL in control animals was 20% to 35% after 4 hours. The polyethylene glycol vehicle was determined to have no effect on CETP activity in this model.

Alternatively, conscious, non-anesthetized animals received an oral gavage dose of test compound as a suspension in 0.1% methyl cellulose in water.
15 At a time determined for each compound at which plasma levels of the test substance reached their peak (C_{max}) after oral dosing, the animals were anesthetized with pentobarbital and then dosed with 0.2 mL of a solution containing [^3H]CE-HDL into the jugular vein as described above. Control animals received 0.25 mL of the vehicle solution without test compound by oral
20 gavage. After 4 hours, the animals were sacrificed, blood samples were collected, and the percentage [^3H]CE transferred from HDL to LDL and VLDL (% transfer) assayed, as described above. The aqueous methyl cellulose vehicle was determined to have no effect on CETP activity in this model. Results from testing in this model are summarized in Table 11.

25 Alternatively, inhibition of CETP activity by a test compound was determined by administering the compound to mice which have been selected for expression of human CETP (hCETP) by transgenic manipulation (hCETP mice). Test compounds were administered by intravenous injection, or oral gavage and the amount of transfer of tritium-labeled cholesteryl ester ([^3H]CE) from HDL to
30 VLDL and LDL particles was determined, and compared to the amount of

transfer observed in control animals. C57Bl/6 mice that were homozygous for the hCETP gene were maintained on a high fat chow diet, such as TD 88051, as described by Nishina et al. (J Lipid Res., 31, 859-869 (1990)) for at least two weeks prior to the study. Mice received an oral gavage dose of test compound as
5 a suspension in 0.1% methyl cellulose in water or an intravenous bolus injection of test compound in 10% ethanol and 90% polyethylene glycol. Control animals received the vehicle solution without test compound by oral gavage or by an intravenous bolus injection. At the start of the experiment all animals received 0.05 mL of a solution containing [3 H]CE-HDL into the tail vein. [3 H]CE-HDL
10 is a preparation of human HDL containing tritium-labeled cholesteryl ester, and was prepared according to the method of Glenn et al. (*Meth. Enzymol.*, 263, 339-351 (1996)). After 30 minutes, the animals were exsanguinated and blood collected in standard microtainer tubes containing ethylenediamine tetraacetic acid. Blood samples were mixed well and kept on ice until the completion of the
15 experiment. Plasma was obtained by centrifugation of the blood samples at 4 °C. The plasma was separated and analyzed by gel filtration chromatography and the relative proportion of [3 H]CE in the VLDL, LDL and HDL regions was determined.

The percentage [3 H]CE transferred from HDL to LDL and VLDL (%
20 transfer) was calculated based on the total radioactivity in equivalent plasma samples before precipitation. Typically, the amount of transfer from HDL to LDL and VLDL in control animals was 20% to 35% after 30 min. The polyethylene glycol and the aqueous methyl cellulose vehicles were determined to have no effect on CETP activity in this model. Results from testing in this
25 model are summarized in Table 12.

ASSAY OF PLASMA HDL ELEVATION *IN VIVO*.

Syrian Golden hamsters were made hypercholesterolemic by feeding cholesterol supplemented chow for a minimum of two weeks, as described above. Test compounds were administered orally in selected aqueous or oil based
30 vehicles for up to 1 week. Serum was obtained and analyzed by precipitation or

size exclusion chromatography for the relative abundance of VLDL, LDL and HDL. Results from testing in this model are summarized in Table 13.

Alternatively, a strain of C57bl mouse was made to transgenically express human CETP. Plasma concentrations of hCETP ranged from 2-20
5 $\mu\text{g/ml}$. The hCETP mice were made hypercholesterolemic by feeding cholesterol and fat supplemented chow for a minimum of two weeks, as described above. Test compounds were administered orally in selected aqueous or oil based vehicles for up to 1 week. Serum was obtained and analyzed by size exclusion chromatography for the relative abundance of
10 VLDL, LDL and HDL. Results from testing in this model are summarized in Table 14.

Alternatively, cynomologous monkeys were maintained on a normal chow diet. The compound corresponding to example 8 was dissolved in a corn oil based vehicle and administered by oral gavage at 10 mpk q.d. for up
15 to 11 days. Plasma levels of drug were detected throughout the experiment in treated animals at ranges of 0.1-1.5 $\mu\text{g/mL}$. Periodically, plasma samples were taken and analyzed for total cholesterol and HDL. After seven days, the treated animals exhibited a 2% increase in HDL and a 5% increase in total cholesterol, relative to vehicle-treated controls.

20 Alternatively, rabbits were maintained on a normal chow diet. The compound corresponding to example 8 was dissolved in a vehicle of ethanol:propylene glycol (1.5:18) and administered by Alzet pump at 30 mg/day/animal for up to 14 days. Plasma concentrations of drug were detected throughout the duration of the pump infusion in treated animals and
25 averaged 1.2 $\mu\text{g/mL}$. Periodically, plasma samples were taken and analyzed for triglycerides, total cholesterol, and HDL. After fourteen days, the treated animals exhibited a 12% decrease in HDL, a 19% decrease in total cholesterol, as well as a 17% increase in triglycerides, compared to pre-dose levels.

Table 9. Inhibition of CETP Activity by Examples in Reconstituted Buffer Assay.

| <u>Ex.</u> <u>No.</u> | <u>IC₅₀</u> (μ M) | <u>Ex.</u> <u>No.</u> | <u>IC₅₀</u> (μ M) |
|--------------------------|--------------------------------------|--------------------------|--------------------------------------|
| 8 | 0.0008 | 42 | 0.38 |
| 11 | 0.001 | 27 | 0.44 |
| 19 | 0.004 | 26 | 0.53 |
| 9 | 0.008 | 29 | 0.72 |
| 10 | 0.012 | 3 | 0.76 |
| 2 | 0.014 | 28 | 0.86 |
| 4 | 0.014 | 32 | 1.2 |
| 20 | 0.027 | 25 | 1.4 |
| 22 | 0.027 | 39 | 1.6 |
| 12 | 0.034 | 15 | 1.6 |
| 14 | 0.04 | 30 | 2.7 |
| 18 | 0.044 | 33B | 3.2 |
| 16 | 0.049 | 5 | 3.4 |
| 43 | 0.058 | 31 | 3.5 |
| 23 | 0.066 | 7 | 4.9 |
| 34 | 0.076 | 44 | 6.8 |
| 41 | 0.086 | 17 | 18 |
| 21 | 0.11 | 6 | 68 |
| 13 | 0.13 | 44A | > 50 |
| 1 | 0.14 | | |
| 33 | 0.15 | | |
| 38 | 0.18 | | |
| 36 | 0.20 | | |
| 37 | 0.21 | | |
| 40 | 0.23 | | |
| 35 | 0.28 | | |
| 24 | 0.33 | | |

Table 10. Inhibition of CETP Activity by
Examples in Human Plasma Assay.

| <u>Ex.</u> <u>No.</u> | <u>IC₅₀</u> <u>(μM)</u> |
|--------------------------|--|
| 8 | 0.049 |
| 11 | 0.072 |
| 10 | 0.11 |
| 22 | 0.14 |
| 19 | 0.19 |
| 20 | 0.3 |
| 18 | 0.44 |
| 14 | 0.59 |
| 9 | 0.62 |
| 2 | 0.65 |
| 4 | 0.65 |
| 16 | 0.77 |
| 12 | 0.79 |
| 34 | 1.4 |
| 43 | 1.5 |
| 23 | 2.0 |
| 1 | 5.6 |
| 41 | 7.2 |
| 42 | 11 |
| 3 | 20 |

Table 11. Inhibition of CETP-mediated Transfer in Hamster

| Ex. No. | Single Oral Dose | % Inhibition of Transfer |
|--------------------|-----------------------------|-------------------------------------|
| 8 | 10 mpk | 35 |

Table 12. Inhibition of CETP-mediated Transfer in hCETP Mice.

| Ex. No. | Single Oral Dose | % Inhibition of Transfer |
|---------|------------------|--------------------------|
| 8 | 60 mpk | 40 |

Table 13. Change in Lipoprotein Profile in Hamster.

| Ex. No. | Oral Dose qd, 5 days | % Change in Lipoprotein Profile | | |
|---------|----------------------|---------------------------------|-----|------|
| | | HDL | LDL | VLDL |
| 8 | 30 mpk | 12 | -12 | -22 |

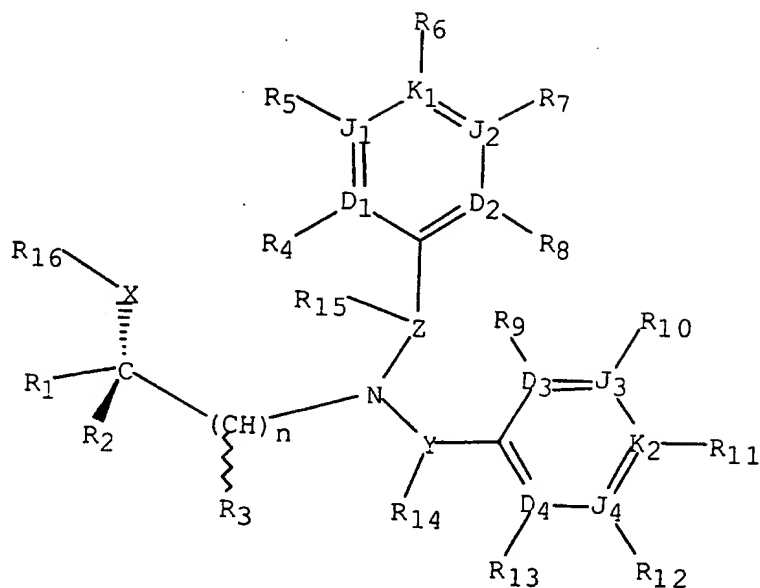
Table 14. Change in Lipoprotein Profile in hCETP Mice.

| Ex. No. | Oral Dose qd, 5 days | % Change in Lipoprotein Profile | | |
|---------|----------------------|---------------------------------|-----|------|
| | | HDL | LDL | VLDL |
| 8 | 30 mpk | 12 | 20 | -- |

212

What we claim is:

1. A compound having the formula:



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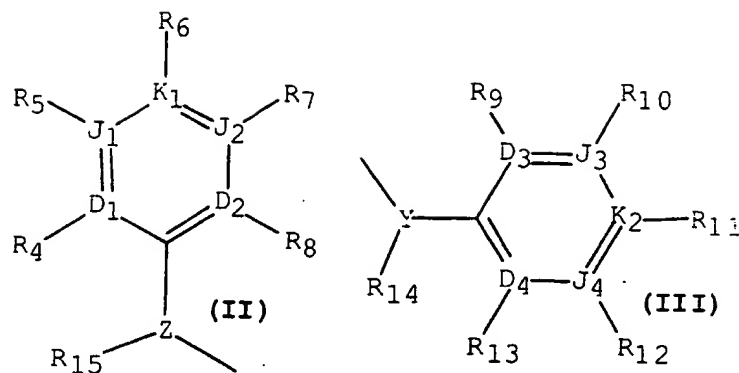
and a pharmaceutically-acceptable salt thereof, wherein;

n is an integer selected from 1 through 4;

X is oxy;

R₁ is selected from the group consisting of haloalkyl, haloalkenyl,

10 haloalkoxymethyl, and haloalkenyloxymethyl with the proviso that R₁ has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R₂ and (CHR₃)_n-N(A)Q wherein A is Formula (II) and Q is Formula (III);



R₁₆ is selected from the group consisting of hydrido, alkyl, acyl, aroyl, heteroaroyl, trialkylsilyl, and a spacer selected from the group consisting of a covalent single bond and a linear spacer moiety having a chain length of 1 to 4 atoms linked to the point of bonding of any aromatic substituent selected from the group consisting of R₄, R₈, R₉, and R₁₃ to form a heterocyclyl ring having from 5 through 10 contiguous members:

D₁, D₂, J₁, J₂ and K₁ are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more than one of D₁, D₂, J₁, J₂ and K₁ is a covalent bond, no more than one of D₁, D₂, J₁, J₂ and K₁ is O, no more than one of D₁, D₂, J₁, J₂ and K₁ is S, one of D₁, D₂, J₁, J₂ and K₁ must be a covalent bond when two of D₁, D₂, J₁, J₂ and K₁ are O and S, and no more than four of D₁, D₂, J₁, J₂ and K₁ are N;

D₃, D₄, J₃, J₄ and K₂ are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more than one is a covalent bond, no more than one of D₃, D₄, J₃, J₄ and K₂ is O, no more than one of D₃, D₄, J₃, J₄ and K₂ is S, no more than two of D₃, D₄,

J_3 , J_4 and K_2 are O and S. one of D_3 , D_4 , J_3 , J_4 and K_2 must be a covalent bond when two of D_3 , D_4 , J_3 , J_4 and K_2 are O and S. and no more than four of D_3 , D_4 , J_3 , J_4 and K_2 are N:

R_2 is selected from the group consisting of hydrido, aryl, aralkyl,

- 5 alkyl, alkenyl, alkenyloxyalkyl, haloalkyl, haloalkenyl, halocycloalkyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, halocycloalkoxy, halocycloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, heteroaryl, dicyanoalkyl, and carboalkoxycyanoalkyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n$ -
- 10 N(A)Q;

R_3 is selected from the group consisting of hydrido, hydroxy, cyano, aryl, aralkyl, acyl, alkoxy, alkyl, alkenyl, alkoxyalkyl, heteroaryl, alkenyloxyalkyl, haloalkyl, haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl, monocyanoalkyl, dicyanoalkyl, carboxamide, and

15 carboxamidoalkyl with the provisos that $(CHR_3)_n$ -N(A)Q has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from a group consisting of a covalent single bond,

$(C(R_{14})_2)_q$ wherein q is an integer selected from 1 and 2 and $(CH(R_{14}))_g$ -W-

- 20 $(CH(R_{14}))_p$ wherein g and p are integers independently selected from 0 and 1;

R_{14} is selected from the group consisting of hydrido, hydroxy, cyano, hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkynyl, alkoxyalkyl, haloalkyl,

215

haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl,

monocarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl,

carboalkoxycyanoalkyl, carboalkoxy, carboxamide, carboxamidoalkyl;

Z is selected from the group consisting of covalent single bond.

- 5 $(C(R_{15})_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH(R_{15}))_j$ -W-
 $(CH(R_{15}))_k$ wherein j and k are integers independently selected from 0 and 1;

W is selected from the group consisting of O, C(O), C(S),

C(O)N(R₁₄), C(S)N(R₁₄), (R₁₄)NC(O), (R₁₄)NC(S), S, S(O), S(O)₂,

- 10 S(O)₂N(R₁₄), (R₁₄)NS(O)₂, and N(R₁₄) with the proviso that R₁₄ is other
 than cyano;

R₁₅ is selected from the group consisting of hydrido, cyano,

hydroxyalkyl, acyl, alkoxy, alkyl, alkenyl, alkynyl, alkoxyalkyl, haloalkyl,

haloalkenyl, haloalkoxy, haloalkoxyalkyl, haloalkenyloxyalkyl,

- 15 monocarboalkoxyalkyl, monocyanoalkyl, dicyanoalkyl,

carboalkoxycyanoalkyl, carboalkoxy, carboxamide, and carboxamidoalkyl;

R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁, R₁₂, and R₁₃ are independently

selected from the group consisting of hydrido, carboxy, heteroaralkylthio,

heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy,

- 20 heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl,

perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl,

aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl,

cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl,

heteroaryl amino, N-heteroaryl amino-N-alkyl amino,

- 25 heteroaryl aminoalkyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl,

haloalkoxyalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy,

cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl,

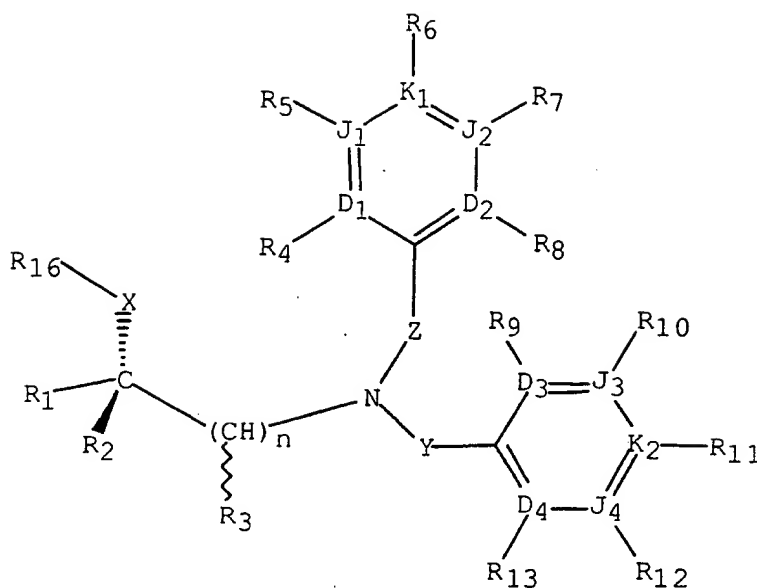
- cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro, lower alkylamino, alkylthio, alkylthioalkyl, arylamino, aralkylamino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, 5 arylsulfinylalkyl, arylsulfonylalkyl, heteroarylsulfinylalkyl, heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, 10 diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, 15 cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, 20 heteroaryloxy, heteroaryloxyalkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and 25 diaralkoxyphosphonoalkyl with the proviso that and R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} , R_{12} , and R_{13} are each independently selected to maintain the tetravalent nature of carbon, trivalent nature of nitrogen, the divalent nature of sulfur, and the divalent nature of oxygen;

R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} are independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no more than one of the group consisting of spacer pairs R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , is used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} is used at the same time;

R_4 and R_9 , R_4 and R_{13} , R_8 and R_9 , and R_8 and R_{13} are independently selected to form a spacer pair wherein said spacer pair is taken together to form a linear moiety wherein said linear moiety forms a ring selected from the group consisting of a partially saturated heterocyclyl ring having from 5 through 8 contiguous members and a heteroaryl ring having from 5 through 6 contiguous members with the proviso that no more than one of the group consisting of spacer pairs R_4 and R_9 , R_4 and R_{13} , R_8 and R_9 , and R_8 and R_{13} is used at the same time.

2. The compound as recited in Claim 1 having the formula of:

218



or a pharmaceutically acceptable salt thereof, wherein;

n is an integer selected from 1 through 3;

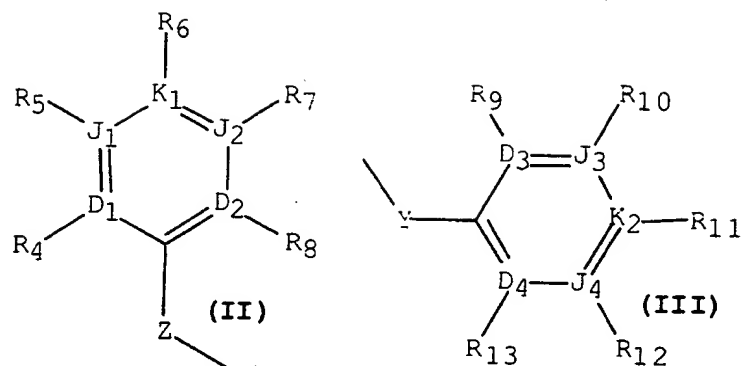
5 X is oxy;

R_{16} is selected from the group consisting of hydrido and a spacer selected from the group consisting of a covalent single bond and a linear spacer moiety having a chain length of 1 to 4 atoms linked to the point of bonding of any aromatic substituent selected from the group consisting of R_4 ,

10 R_8 , R_9 , and R_{13} to form a heterocyclyl ring having from 5 through 10 contiguous members;

R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);

15



- D₁, D₂, J₁, J₂ and K₁ are independently selected from the group consisting of C, N, O, S and a covalent bond with the provisos that no more than one of D₁, D₂, J₁, J₂ and K₁ is a covalent bond, no more than one of D₁, D₂, J₁, J₂ and K₁ is O, no more than one of D₁, D₂, J₁, J₂ and K₁ is S, one of D₁, D₂, J₁, J₂ and K₁ must be a covalent bond when two of D₁, D₂, J₁, J₂ and K₁ are O and S, and no more than four of D₁, D₂, J₁, J₂ and K₁ are N;

- D₃, D₄, J₃, J₄ and K₂ are independently selected from the group consisting of C, N, O, S and a covalent bond with the provisos that no more than one of D₃, D₄, J₃, J₄ and K₂ is a covalent bond, no more than one of D₃, D₄, J₃, J₄ and K₂ is O, no more than one of D₃, D₄, J₃, J₄ and K₂ is S, one of D₃, D₄, J₃, J₄ and K₂ must be a covalent bond when two of D₃, D₄, J₃, J₄ and K₂ are O and S, and no more than four of D₃, D₄, J₃, J₄ and K₂ are N;

- R₂ is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, haloalkoxy, haloalkoxyalkyl, perhaloaryl, perhaloaralkyl,

perhaloaryloxyalkyl, and heteroaryl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$:

R_3 is selected from the group consisting of hydrido, aryl, alkyl,

alkenyl, haloalkyl, and haloalkoxyalkyl with the provisos that $(CHR_3)_n-$

- 5 $N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 :

Y is selected from the group consisting of a covalent single bond,

$(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$

wherein j and k are integers independently selected from 0 and 1;

- 10 Z is selected from the group consisting of covalent single bond,

$(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$

wherein j and k are integers independently selected from 0 and 1;

R_4, R_8, R_9 , and R_{13} are independently selected from the group consisting of hydrido, halo, haloalkyl, and alkyl ;

- 15 $R_5, R_6, R_7, R_{10}, R_{11}$, and R_{12} are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl,
- 20 halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroarylamino, N-heteroarylamino-N-alkylamino, heteroarylaminoalkyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxylalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy,
- 25 cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy,

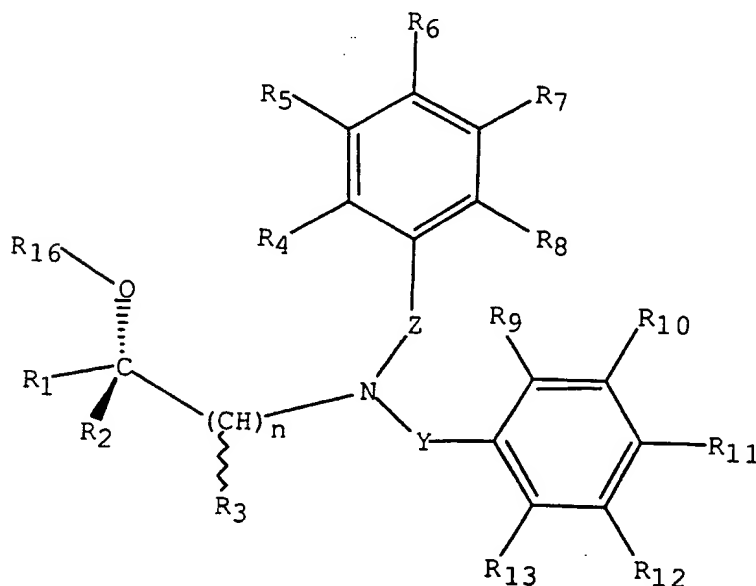
- halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, hydroxy, amino, thio, nitro, lower alkylamino, alkylthio, alkylthioalkyl, arylamino, aralkylamino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, 5 heteroarylsulfinylalkyl, heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, 10 arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower 15 cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, arylalkenyl, 20 heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl;
- 25 R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} are independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a

cycloalkenyl ring having 5 through 8 contiguous members, a partially saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no more than one of the group consisting of spacer pairs R_4 and

- 5 R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , is used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} is used at the same time.

3. The compound as recited in Claim 2 having the formula of:

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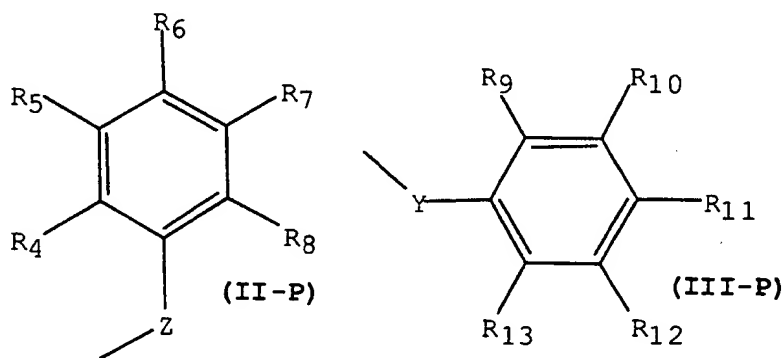
or a pharmaceutically acceptable salt thereof, wherein;

n is an integer selected from 1 through 3;

R_{16} is selected from the group consisting of hydrido and a spacer selected from the group consisting of a covalent single bond and a linear spacer moiety having a chain length of 1 to 4 atoms linked to the point of bonding of any aromatic substituent selected from the group consisting of R_4 .

- 5 R_8 , R_9 , and R_{13} to form a heterocyclyl ring having from 5 through 10 contiguous members;

R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein
 10 Ap is Formula (II-P) and Qp is Formula (III-P):



- R_2 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, haloalkoxy, haloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, and heteroaryl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;
- 15

R_3 is selected from the group consisting of hydrido, aryl, alkyl,

alkenyl, haloalkyl, and haloalkoxyalkyl with the provisos that $(CHR_3)_n$ -
N(Ap)Qp has a lower Cahn-Ingold-Prelog stereochemical system ranking
than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than

5 R_2 ;

Y is selected from the group consisting of a covalent single bond,

$(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j$ -O- $(CH_2)_k$
wherein j and k are integers independently selected from 0 and 1;

Z is selected from the group consisting of covalent single bond,

10 $(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j$ -O- $(CH_2)_k$
wherein j and k are integers independently selected from 0 and 1;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group
consisting of hydrido, halo, haloalkyl, and alkyl ;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the
15 group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy,
cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocycloxy,
aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl,
aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl,
halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl,
20 cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroaryl amino, N-
heteroaryl amino-N-alkyl amino, heteroaryl aminoalkyl, haloalkylthio,
alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxylalkyl, heteroaralkoxy,
cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy,
cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy,
25 halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl,

- hydroxy, amino, thio, nitro, lower alkylamino, alkylthio, alkylthioalkyl, arylamino, aralkylamino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, heteroarylsulfinylalkyl, heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy,
- hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido,
- alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl;

R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and

- R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} are independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially

saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no more than one of the group consisting of spacer pairs R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , is used at the same time and that

5 no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} is used at the same time.

4. The compound as recited in Claim 3 or a pharmaceutically acceptable salt thereof, wherein:

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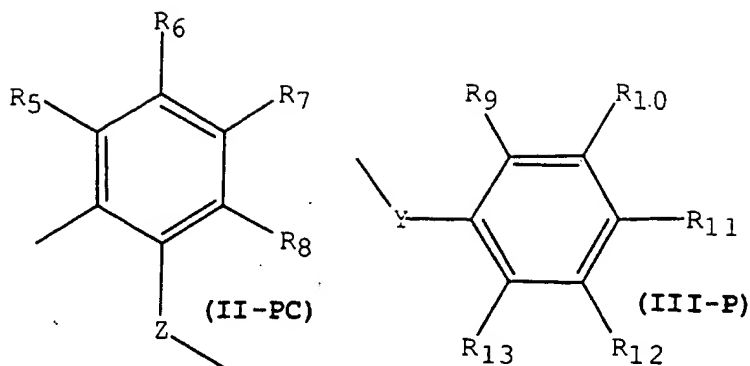
n is the integer 1;

R_{16} is taken together with R_4 , R_8 , R_9 , or R_{13} to form a spacer selected from the group consisting of a covalent single bond, CH_2 , $CH(CH_3)$, CF_2 , $C(O)$, $C(S)$, and SO_2 ;

15

R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Apc)Qp$ wherein Apc is Formula (II-PC) and Qp is Formula (III-

20 P);



R_2 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, vinyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl
 5 with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Apc)Qp$:

R_3 is selected from the group consisting of hydrido, methyl, ethyl, vinyl, phenyl, 4-trifluoromethylphenyl, trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and
 10 pentafluoroethyl with the provisos that $(CHR_3)_n-N(Apc)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from the group consisting of covalent single bond, oxy, methyleneoxy, methylene, and ethylene;

15 Z is selected from the group consisting of covalent single bond, oxy, methyleneoxy, methylene, and ethylene;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

- R_5 and R_{10} are independently selected from the group consisting of 4-aminophenoxy, benzoyl, benzyl, benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-bromo-2-nitrophenoxy, 3-bromobenzyloxy, 4-bromobenzyloxy, 4-bromophenoxy, 5-bromopyrid-2-yloxy,
- 5 4-butoxyphenoxy, chloro, 3-chlorobenzyl, 2-chlorophenoxy, 4-chlorophenoxy, 4-chloro-3-ethylphenoxy, 3-chloro-4-fluorobenzyl, 3-chloro-4-fluorophenyl, 3-chloro-2-fluorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy,
- 10 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy, 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy, 2-cyanopyrid-3-yloxy, 4-cyanophenoxy, cyclobutoxy, cyclobutyl, cyclohexoxy, cyclohexylmethoxy, cyclopentoxy, cyclopentyl, cyclopentylcarbonyl, cyclopropyl, cyclopropylmethoxy, cyclopropoxy,
- 15 2,3-dichlorophenoxy, 2,4-dichlorophenoxy, 2,4-dichlorophenyl, 3,5-dichlorophenyl, 3,5-dichlorobenzyl, 3,4-dichlorophenoxy, 3,4-difluorophenoxy, 2,3-difluorobenzyloxy, 2,4-difluorobenzyloxy, 3,4-difluorobenzyloxy, 2,5-difluorobenzyloxy, 3,5-difluorophenoxy, 3,4-difluorophenyl, 3,5-difluorobenzyloxy, 4-difluoromethoxybenzyloxy,
- 20 2,3-difluorophenoxy, 2,4-difluorophenoxy, 2,5-difluorophenoxy, 3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3,4-dimethylbenzyl, 3,4-dimethylbenzyloxy, 3,5-dimethylbenzyloxy, 2,2-dimethylpropoxy, 1,3-dioxan-2-yl, 1,4-dioxan-2-yl, 1,3-dioxolan-2-yl, ethoxy, 4-ethoxyphenoxy,
- 25 4-ethylbenzyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy, 3-ethyl-5-methylphenoxy, fluoro, 4-fluoro-3-methylbenzyl, 4-fluoro-3-methylphenyl, 4-fluoro-3-methylbenzoyl, 4-fluorobenzyloxy, 2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy, 3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzyloxy,

- 3-fluoro-5-trifluoromethylbenzyloxy, 4-fluoro-2-trifluoromethylbenzyloxy,
4-fluoro-3-trifluoromethylbenzyloxy, 2-fluorophenoxy, 4-fluorophenoxy,
2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzyloxy,
4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy,
5 4-fluoropyrid-2-yloxy, 2-furyl, 3-furyl, heptafluoropropyl,
1,1,1,3,3,3-hexafluoropropyl, 2-hydroxy-3,3,3-trifluoropropoxy,
3-iodobenzyloxy, isobutyl, isobutylamino, isobutoxy, 3-isoxazolyl,
4-isoxazolyl, 5-isoxazolyl, isopropoxy, isopropyl, 4-isopropylbenzyloxy,
3-isopropylphenoxy, 4-isopropylphenoxy, isopropylthio,
10 4-isopropyl-3-methylphenoxy, 3-isothiazolyl, 4-isothiazolyl,
5-isothiazolyl, 3-methoxybenzyl, 4-methoxycarbonylbutoxy,
3-methoxycarbonylprop-2-enyloxy, 4-methoxyphenyl,
3-methoxyphenylamino, 4-methoxyphenylamino, 3-methylbenzyloxy,
4-methylbenzyloxy, 3-methylphenoxy, 3-methyl-4-methylthiophenoxy,
15 4-methylphenoxy, 1-methylpropoxy, 2-methylpyrid-5-yloxy,
4-methylthiophenoxy, 2-naphthyloxy, 2-nitrophenoxy, 4-nitrophenoxy,
3-nitrophenyl, 4-nitrophenylthio, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl,
pentafluoroethyl, pentafluoroethylthio, 2,2,3,3,3-pentafluoropropyl,
1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl, phenoxy,
20 phenylamino, 1-phenylethoxy, phenylsulfonyl, 4-propanoylphenoxy,
propoxy, 4-propylphenoxy, 4-propoxyphenoxy, thiophen-3-yl, *sec*-butyl,
4-*sec*-butylphenoxy, *tert*-butoxy, 3-*tert*-butylphenoxy, 4-*tert*-butylphenoxy,
1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
2-(5,6,7,8-tetrahydronaphthyloxy), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl,
25 thiophen-2-yl, 2,3,5-trifluorobenzyloxy, 2,2,2-trifluoroethoxy,
2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
3-trifluoromethoxyphenoxy, 4-trifluoromethoxyphenoxy, trifluoromethyl,
3-trifluoromethylbenzyloxy, 4-trifluoromethylbenzyloxy,
30 2,4-bis-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,

- 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
 3-trifluoromethylphenyl, 3-trifluoromethylthiobenzyloxy,
 4-trifluoromethylthiobenzyloxy, 2,3,4-trifluorophenoxy,
 5 2,3,4-trifluorophenyl, 2,3,5-trifluorophenoxy, 3,4,5-trimethylphenoxy,
 3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy,
 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, and
 trifluoromethylthio;

- R_6 and R_{11} are independently selected from the group consisting of
 10 chloro, fluoro, hydrido, difluoromethoxy, trifluoromethyl, trifluoromethoxy,
 pentafluoroethyl, and 1,1,2,2-tetrafluoroethoxy;

R_7 and R_{12} are independently selected from the group consisting of
 hydrido, fluoro, and trifluoromethyl.

15

5. The compound as recited in Claim 4 or a pharmaceutically acceptable salt
 thereof, wherein:

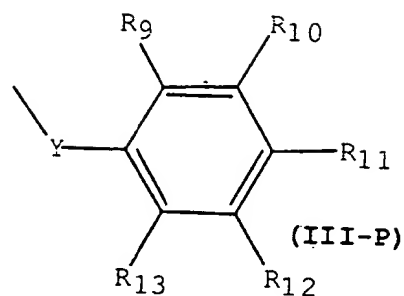
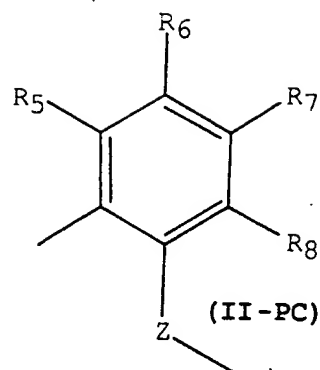
R_{16} is taken together with R_4 , R_8 , R_9 , or R_{13} to form a covalent single
 bond.

20

n is the integer 1;

R_1 is selected from the group consisting of trifluoromethyl and
 pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog
 stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Apc)Qp$ wherein
 Apc is Formula (II-PC) and Qp is Formula (III-P);

231



R_2 is selected from the group consisting of hydrido and phenyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Apc)Qp$:

5 R_3 is selected from the group consisting of hydrido, methyl, trifluoromethyl, and difluoromethyl with the provisos that $(CHR_3)_n-N(Apc)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

10 Y is methylene;
 Z is covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

15 R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy, 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy,

- 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyl)oxy, 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R_{10} is selected from the group consisting of cyclopentyl, 1,1,2,2-tetrafluoroethoxy, 2-furyl, 1,1-bis-trifluoromethyl-1-hydroxymethyl, isobutyl, isopropoxy, pentafluoroethyl, trifluoromethoxy, trifluoromethyl, and trifluoromethylthio;

- 10 R_6 and R_{11} are independently selected from the group consisting of fluoro and hydrido;

R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

15

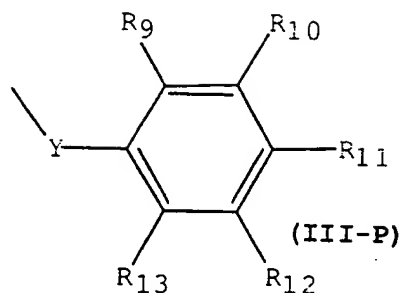
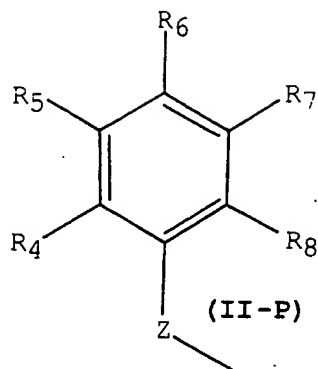
6. The compound as recited in Claim 3 or a pharmaceutically acceptable salt thereof, wherein:

n is an integer selected from 1 and 2;

20

R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);

233



R_{16} is hydrido;

R_2 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, haloalkoxy, haloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, and heteroaryl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:

R_3 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, and haloalkoxyalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from the group consisting of a covalent single bond, oxy and C1-C2 alkylene;

Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

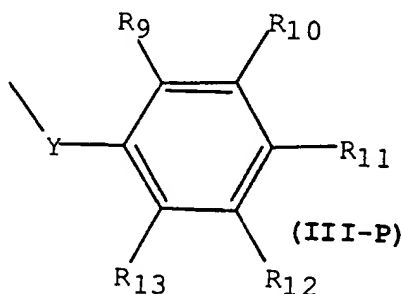
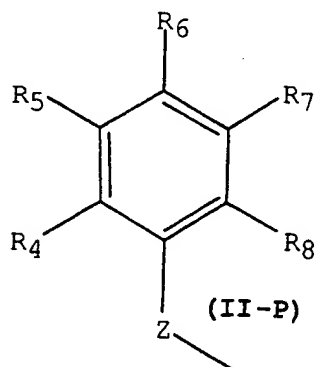
- R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

- 10 7. The compound as recited in Claim 6 or a pharmaceutically acceptable salt thereof, wherein;

n is the integer 1;

R_{16} is hydrido;

- R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n$ -N(Ap)Qp wherein Ap is Formula (II-P) and Qp is Formula (III-P);



;

R_2 is selected from the group consisting of hydrido, methyl, ethyl, propyl, butyl, vinyl, phenyl, 4-trifluoromethylphenyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-

- 5 Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:

R_3 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, methyl, ethyl, vinyl, trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-

- 10 Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-

Ingold-Prelog stereochemical system ranking than R_2 :

Y is selected from the group consisting of methylene and ethylene:

Z is a covalent single bond;

- R_4 , R_8 , R_9 , and R_{13} are independently selected from the group
- 15 consisting of hydrido and fluoro;

- R_5 and R_{10} are independently selected from the group consisting of 4-aminophenoxy, benzoyl, benzyl, benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-bromo-2-nitrophenoxy, 3-bromobenzyloxy, 4-bromobenzyloxy, 4-bromophenoxy, 5-bromopyrid-2-yloxy,
- 20 4-butoxyphenoxy, chloro, 3-chlorobenzyl, 2-chlorophenoxy, 4-chlorophenoxy, 4-chloro-3-ethylphenoxy, 3-chloro-4-fluorobenzyl, 3-chloro-4-fluorophenyl, 3-chloro-2-fluorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy,

- 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy,
4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy,
2-cyanopyrid-3-yloxy, 4-cyanophenoxy, cyclobutoxy, cyclobutyl,
cyclohexoxy, cyclohexylmethoxy, cyclopentoxy, cyclopentyl,
5 cyclopentylcarbonyl, cyclopropyl, cyclopropylmethoxy, cyclopropoxy,
2,3-dichlorophenoxy, 2,4-dichlorophenoxy, 2,4-dichlorophenyl,
3,5-dichlorophenyl, 3,5-dichlorobenzyl, 3,4-dichlorophenoxy,
3,4-difluorophenoxy, 2,3-difluorobenzoyloxy, 2,4-difluorobenzoyloxy,
3,4-difluorobenzoyloxy, 2,5-difluorobenzoyloxy, 3,5-difluorophenoxy,
10 3,4-difluorophenyl, 3,5-difluorobenzoyloxy, 4-difluoromethoxybenzoyloxy,
2,3-difluorophenoxy, 2,4-difluorophenoxy, 2,5-difluorophenoxy,
3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy, 3,5-dimethylphenoxy,
3,4-dimethylphenoxy, 3,4-dimethylbenzyl, 3,4-dimethylbenzoyloxy,
3,5-dimethylbenzoyloxy, 2,2-dimethylpropoxy, 1,3-dioxan-2-yl,
15 1,4-dioxan-2-yl, 1,3-dioxolan-2-yl, ethoxy, 4-ethoxyphenoxy,
4-ethylbenzoyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy,
3-ethyl-5-methylphenoxy, fluoro, 4-fluoro-3-methylbenzyl,
4-fluoro-3-methylphenyl, 4-fluoro-3-methylbenzoyl, 4-fluorobenzoyloxy,
2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy,
20 3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzoyloxy,
3-fluoro-5-trifluoromethylbenzoyloxy, 4-fluoro-2-trifluoromethylbenzoyloxy,
4-fluoro-3-trifluoromethylbenzoyloxy, 2-fluorophenoxy, 4-fluorophenoxy,
2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzoyloxy,
4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy,
25 4-fluoropyrid-2-yloxy, 2-furyl, 3-furyl, heptafluoropropyl,
1,1,1,3,3,3-hexafluoropropyl, 2-hydroxy-3,3,3-trifluoropropoxy,
3-iodobenzoyloxy, isobutyl, isobutylamino, isobutoxy, 3-isoxazolyl,
4-isoxazolyl, 5-isoxazolyl, isopropoxy, isopropyl, 4-isopropylbenzoyloxy,
3-isopropylphenoxy, 4-isopropylphenoxy, isopropylthio,
30 4-isopropyl-3-methylphenoxy, 3-isothiazolyl, 4-isothiazolyl,

- 5-isothiazolyl, 3-methoxybenzyl, 4-methoxycarbonylbutoxy,
3-methoxycarbonylprop-2-enyloxy, 4-methoxyphenyl,
3-methoxyphenylamino, 4-methoxyphenylamino, 3-methylbenzyloxy,
4-methylbenzyloxy, 3-methylphenoxy, 3-methyl-4-methylthiophenoxy,
5 4-methylphenoxy, 1-methylpropoxy, 2-methylpyrid-5-yloxy,
4-methylthiophenoxy, 2-naphthyloxy, 2-nitrophenoxy, 4-nitrophenoxy,
3-nitrophenyl, 4-nitrophenylthio, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl,
pentafluoroethyl, pentafluoroethylthio, 2,2,3,3,3-pentafluoropropyl,
1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl, phenoxy,
10 phenylamino, 1-phenylethoxy, phenylsulfonyl, 4-propanoylphenoxy,
propoxy, 4-propylphenoxy, 4-propoxyphenoxy, thiophen-3-yl, *sec*-butyl,
4-*sec*-butylphenoxy, *tert*-butoxy, 3-*tert*-butylphenoxy, 4-*tert*-butylphenoxy,
1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
2-(5,6,7,8-tetrahydronaphthyloxy), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl,
15 thiophen-2-yl, 2,3,5-trifluorobenzyloxy, 2,2,2-trifluoroethoxy,
2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
3-trifluoromethoxyphenoxy, 4-trifluoromethoxyphenoxy, trifluoromethyl,
3-trifluoromethylbenzyloxy, 4-trifluoromethylbenzyloxy,
20 2,4-bis-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,
3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
3-trifluoromethylphenyl, 3-trifluoromethylthiobenzyloxy,
4-trifluoromethylthiobenzyloxy, 2,3,4-trifluorophenoxy,
25 2,3,4-trifluorophenyl, 2,3,5-trifluorophenoxy, 3,4,5-trimethylphenoxy,
3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy,
3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, and
trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy, trifluoromethyl, and trifluoromethoxy;

R_7 and R_{12} are independently selected from the group consisting of
5 hydrido, fluoro, and trifluoromethyl.

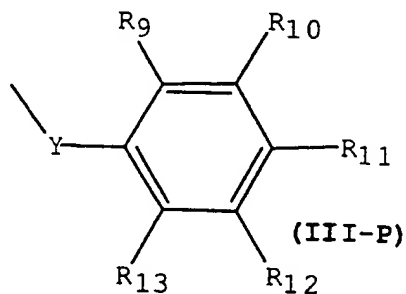
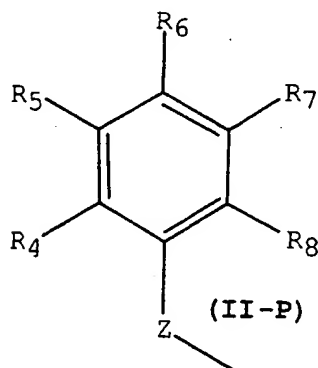
8. The compound as recited in Claim 7 or a pharmaceutically acceptable salt thereof, wherein;

10

n is the integer 1;

R_{16} is hydrido;

R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl,
15 chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



- R_2 is selected from the group consisting of hydrido, methyl, ethyl, phenyl, 4-trifluoromethylphenyl, trifluoromethoxymethyl, 1,1,2,2-tetrafluoroethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:

- R_3 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, methyl, trifluoromethyl, difluoromethyl, and chlorodifluoromethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 :

Y is methylene;

Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

- R_5 and R_{10} are independently selected from the group consisting of benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 3-bromobenzyloxy, 4-bromophenoxy, 4-butoxyphenoxy, 3-chlorobenzyloxy, 2-chlorophenoxy, 4-chloro-3-ethylphenoxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy, 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy, 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy, cyclobutoxy, cyclobutyl, cyclohexylmethoxy, cyclopentoxo, cyclopentyl, cyclopentylcarbonyl, cyclopropylmethoxy, 2,3-dichlorophenoxy,

- 2,4-dichlorophenoxy, 2,4-dichlorophenyl, 3,5-dichlorophenyl,
3,5-dichlorobenzyl, 3,4-dichlorophenoxy, 3,4-difluorophenoxy,
2,3-difluorobenzyl, 3,5-difluorobenzyl, difluoromethoxy,
3,5-difluorophenoxy, 3,4-difluorophenyl, 2,3-difluorophenoxy,
5 2,4-difluorophenoxy, 2,5-difluorophenoxy, 3,5-dimethoxyphenoxy,
3-dimethylaminophenoxy, 3,4-dimethylbenzyl, 3,5-dimethylbenzyl,
3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 1,3-dioxolan-2-yl,
3-ethylbenzyl, 3-ethylphenoxy, 4-ethylaminophenoxy,
3-ethyl-5-methylphenoxy, 4-fluoro-3-methylbenzyl, 4-fluorobenzyl,
10 2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy,
3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzyl,
3-fluoro-5-trifluoromethylbenzyl, 2-fluorophenoxy, 4-fluorophenoxy,
2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzyl,
4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy, 2-furyl, 3-furyl,
15 heptafluoropropyl, 1,1,1,3,3,3-hexafluoropropyl,
2-hydroxy-3,3,3-trifluoropropoxy, isobutoxy, isobutyl, 3-isoxazolyl,
4-isoxazolyl, 5-isoxazolyl, isopropoxy,
3-isopropylbenzyl, 3-isopropylphenoxy, isopropylthio,
4-isopropyl-3-methylphenoxy, 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl,
20 3-methoxybenzyl, 4-methoxyphenylamino, 3-methylbenzyl,
4-methylbenzyl, 3-methylphenoxy, 3-methyl-4-methylthiophenoxy,
4-methylphenoxy, 1-methylpropoxy, 2-methylpyrid-5-yl,
4-methylthiophenoxy, 2-naphthyl, 2-nitrophenoxy, 4-nitrophenoxy,
3-nitrophenyl, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl, pentafluoroethyl,
25 pentafluoroethylthio, 2,2,3,3,3-pentafluoropropyl,
1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl, phenoxy,
phenylamino, 1-phenylethoxy, 4-propylphenoxy, 4-propoxyphenoxy,
thiophen-3-yl, tert-butyl, 3-tert-butylphenoxy, 4-tert-butylphenoxy,
1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,

- 2-(5.6.7.8-tetrahydronaphthoxy), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl, thiophen-2-yl, 2.2.2-trifluoroethoxy, 2.2.2-trifluoroethyl, 3.3.3-trifluoro-2-hydroxypropyl, trifluoromethoxy, 3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy, 4-trifluoromethoxyphenoxy, 3-trifluoromethoxyphenoxy, trifluoromethyl, 3-trifluoromethylbenzyloxy, 1.1-bis-trifluoromethyl-1-hydroxymethyl, 3-trifluoromethylbenzyl, 3.5-bis-trifluoromethylbenzyloxy, 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy, 3-trifluoromethylphenyl, 2,3,4-trifluorophenoxy, 2.3.5-trifluorophenoxy, 3.4.5-trimethylphenoxy, 3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy, 3-(1.1.2.2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, 3-trifluoromethylthiobenzyloxy, and trifluoromethylthio;

- R_6 and R_{11} are independently selected from the group consisting of chloro, fluoro, hydrido, pentafluoroethyl, 1.1.2.2-tetrafluoroethoxy, and trifluoromethyl;

R_7 and R_{12} are independently selected from the group consisting of hydrido, fluoro, and trifluoromethyl.

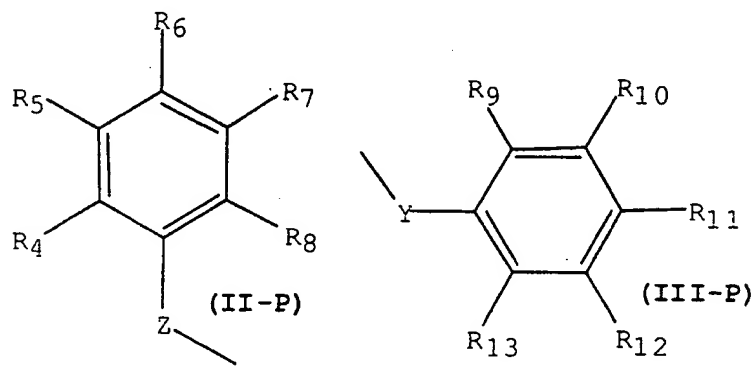
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9. The compound as recited in Claim 6 or a pharmaceutically acceptable salt, wherein;

25 n is the integer 1;

R_{16} is hydrido;

R_1 is haloalkyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P):



R_2 is selected from the group consisting of hydrido, alkyl, haloalkyl, aryl, and haloalkoxy with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$;

10 R_3 is selected from the group consisting of hydrido, alkyl, and haloalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is a covalent single bond;

15 Z is covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

10 10. The compound as recited in Claim 9 or a pharmaceutically acceptable salt thereof, wherein:

n is the integer 1;

R_1 is selected from the group consisting of trifluoromethyl, chlorodifluoromethyl, and pentafluoroethyl:

R_{16} is hydrido;

R_2 is hydrido;

R_3 is hydrido;

Y is a covalent single bond;

20 Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy, 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy,

- 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyl)oxy, 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R_{10} is selected from the group consisting of 1,1,2,2-tetrafluoroethoxy, pentafluoroethyl, and trifluoromethyl;

R_6 and R_{11} are independently selected from the group consisting of fluoro and hydrido;

- 10 R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

11. The compound as recited in Claim 6 or a pharmaceutically acceptable salt, wherein;

15

n is the integer 1;

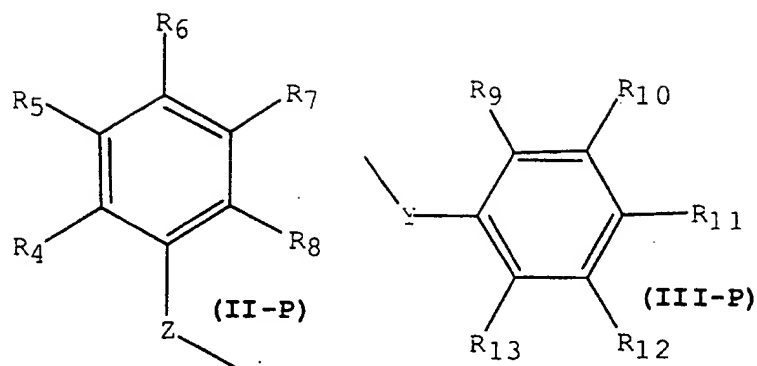
R_{16} is hydrido;

R_1 is haloalkyl with the proviso that R_1 has a higher Cahn-Ingold-

Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$.

- 20 wherein Ap is Formula (II-P) and Qp is Formula (III-P);

245



R₂ is selected from the group consisting of hydrido, alkyl, haloalkyl, aryl, and haloalkoxy with the proviso that R₂ has a lower Cahn-Ingold-Prelog system ranking than both R₁ and (CHR₃)_n-N(Ap)Qp:

R₃ is selected from the group consisting of hydrido, alkyl, and haloalkyl with the provisos that (CHR₃)_n-N(Ap)Qp has a lower Cahn-Ingold-Prelog stereochemical system ranking than R₁ and a higher Cahn-Ingold-Prelog stereochemical system ranking than R₂:

10 Y is oxy;

Z is a covalent single bond;

R₄, R₈, R₉, and R₁₃ are independently selected from the group consisting of hydrido and halo;

15 R₅, R₆, R₇, R₁₀, R₁₁, and R₁₂ are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl,

heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl,
heteroarylthio, and heteroarylsulfonyl.

- 5 12. The compound as recited in Claim 11 or a pharmaceutically acceptable salt thereof, wherein:

n is the integer 1;

R₁ is selected from the group consisting of trifluoromethyl,

- 10 chlorodifluoromethyl, and pentafluoroethyl;

R₁₆ is hydrido;

R₂ is hydrido;

R₃ is hydrido;

Y is oxy;

- 15 Z is a covalent single bond;

R₄, R₈, R₉, and R₁₃ are independently selected from the group consisting of hydrido and fluoro;

- R₅ is selected from the group consisting of 5-bromo-2-fluorophenoxy, 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-
20 difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy, 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyl)oxy),
25 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R_{10} is selected from the group consisting of 1,1,2,2-tetrafluoroethoxy, pentafluoroethyl, and trifluoromethyl;

R_6 and R_{11} are independently selected from the group consisting of fluoro and hydrido;

5 R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

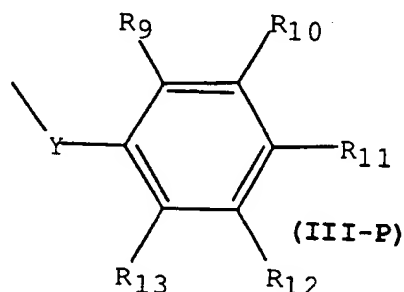
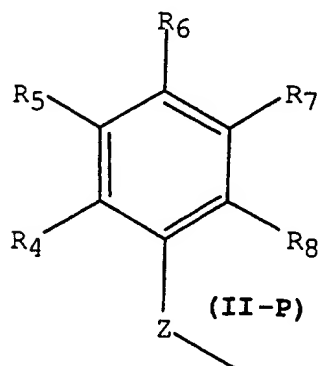
13. The compound as recited in Claim 6 or a pharmaceutically acceptable
10 salt, wherein:

n is the integer 1;

R_{16} is hydrido;

R_1 is haloalkyl with the proviso that R_1 has a higher Cahn-Ingold-

15 Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P);



R_2 is selected from the group consisting of hydrido, alkyl, haloalkyl, aryl, and haloalkoxy with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:

R_3 is selected from the group consisting of hydrido, alkyl, and haloalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 :

Y is methylene;

Z is a covalent single bond;

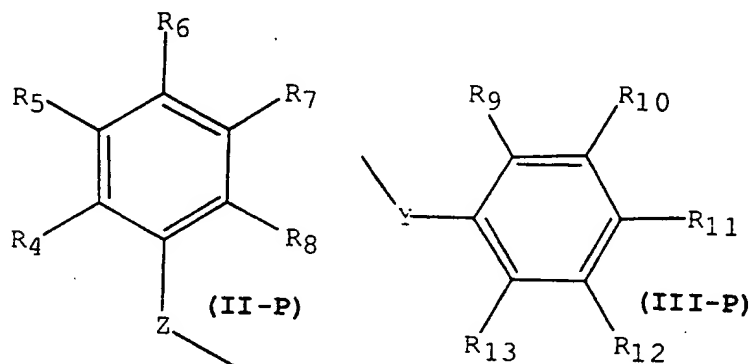
$R_4, R_8, R_9,$ and R_{13} are independently selected from the group consisting of hydrido and halo;

$R_5, R_6, R_7, R_{10}, R_{11},$ and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

14. The compound as recited in Claim 13 or a pharmaceutically acceptable salt thereof, wherein;

n is the integer 1;

R_1 is selected from the group consisting of trifluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(Ap)Qp$ wherein Ap is Formula (II-P) and Qp is Formula (III-P):



5

R_{16} is hydrido;

R_2 is selected from the group consisting of hydrido and phenyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both

10 R_1 and $(CHR_3)_n-N(Ap)Qp$;

R_3 is selected from the group consisting of hydrido, methyl, trifluoromethyl, and difluoromethyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

15

Y is methylene;

Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy, 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy, 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyl)oxy, 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R_{10} is selected from the group consisting of cyclopentyl, 1,1,2,2-tetrafluoroethoxy, 2-furyl, 1,1-bis-trifluoromethyl-1-hydroxymethyl, isobutyl, isopropoxy, pentafluoroethyl, trifluoromethoxy, trifluoromethyl, and trifluoromethylthio;

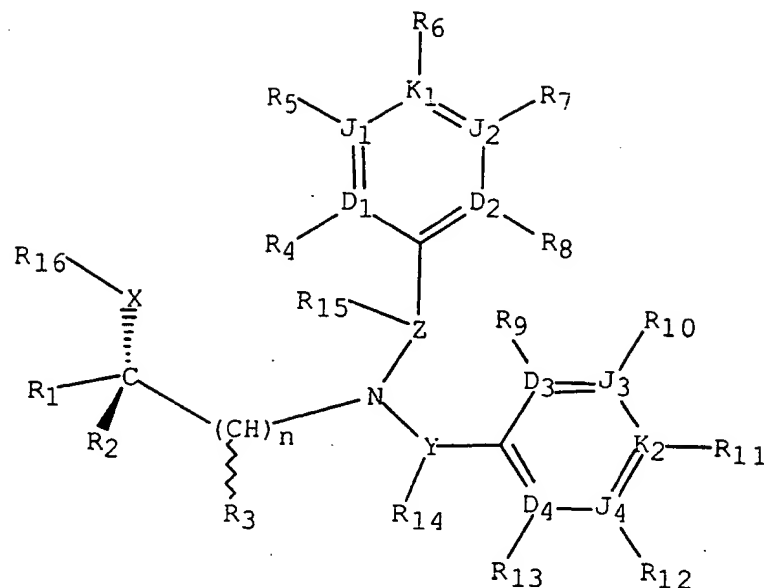
R_6 and R_{11} are independently selected from the group consisting of fluoro and hydrido;

R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

20

15. A compound as recited in Claim 2 having the formula:

251

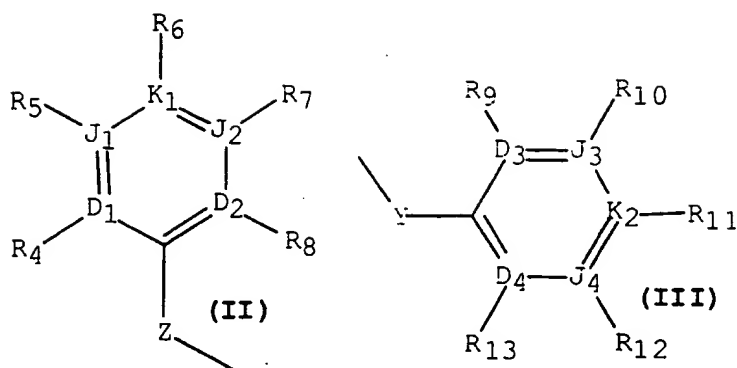


or a pharmaceutically acceptable salt thereof, wherein:

- D_1, D_2, J_1, J_2 and K_1 are each carbon with the proviso that at least one
- 5 of D_3, D_4, J_3, J_4 and K_2 is selected from the group consisting of O, S, and N,
- wherein D_3, D_4, J_3, J_4 and K_2 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more than one of D_3, D_4, J_3, J_4 and K_2 is a covalent bond, no more than one of D_3, D_4, J_3, J_4 and K_2 is O, no more than one of D_3, D_4, J_3, J_4 and K_2 is S, one of
- 10 D_3, D_4, J_3, J_4 and K_2 must be a covalent bond when two of D_3, D_4, J_3, J_4 and K_2 are O and S, and no more than four of D_3, D_4, J_3, J_4 and K_2 are N;

D_1, D_2, J_1, J_2 and K_1 are independently selected from the group consisting of C, O, S, N and covalent bond with the provisos that D_3, D_4, J_3, J_4 and K_2 are each carbon and at least one of D_1, D_2, J_1, J_2 and K_1 is selected

- from the group consisting of O, S, and N wherein, when D_1 , D_2 , J_1 , J_2 and K_1 are selected from the group consisting of C, O, S, covalent bond, and N, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 is a covalent bond, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 is O, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 is S.
- 5 one of D_1 , D_2 , J_1 , J_2 and K_1 must be a covalent bond when two of D_1 , D_2 , J_1 , J_2 and K_1 are O and S, and no more than four of D_1 , D_2 , J_1 , J_2 and K_1 are N;
 n is an integer selected from 1 and 2;
 X is oxy;
- R_{16} is selected from the group consisting of hydrido and a spacer
- 10 selected from the group consisting of a covalent single bond and a linear spacer moiety having a chain length of 1 to 4 atoms linked to the point of bonding of any aromatic substituent selected from the group consisting of R_4 , R_8 , R_9 , and R_{13} to form a heterocyclyl ring having from 5 through 10 contiguous members;
- 15 R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);



R_2 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, haloalkoxy, haloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, and heteroaryl with the proviso that R_2 has a lower

- 5 Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(Ap)Qp$:

R_3 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, and haloalkoxyalkyl with the provisos that $(CHR_3)_n-N(Ap)Qp$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than

- 10 R_2 ;

Y is selected from the group consisting of a covalent single bond, $(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$ wherein j and k are integers independently selected from 0 and 1;

Z is selected from the group consisting of covalent single bond;

- 15 $(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$ wherein j and k are integers independently selected from 0 and 1;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido, halo, haloalkyl, and alkyl :

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, 5 aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroaryl amino, N- 10 heteroaryl amino-N-alkyl amino, heteroaryl aminoalkyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxyalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, 15 hydroxy, amino, thio, nitro, lower alkyl amino, alkylthio, alkylthioalkyl, aryl amino, aralkyl amino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, heteroaryl sulfinylalkyl, heteroaryl sulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, 20 alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoaryl amidosulfonyl, arylsulfonamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroaryl sulfinyl, heteroaryl sulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, 25 aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl,

haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl;

R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and

- 10 R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} are independently selected to form spacer pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially
- 15 saturated heterocyclyl ring having 5 through 8 contiguous members, a heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no more than one of the group consisting of spacer pairs R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , is used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and
- 20 R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} is used at the same time.

16. The compound as recited in Claim 15 or a pharmaceutically acceptable salt thereof, wherein;

D_1, D_2, J_1, J_2 and K_1 are each carbon with the proviso that at least one of D_3, D_4, J_3, J_4 and K_2 is selected from the group consisting of O, S, and N, wherein D_3, D_4, J_3, J_4 and K_2 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more than one of D_3, D_4, J_3, J_4 and K_2 is a covalent bond, no more than one of D_3, D_4, J_3, J_4 and K_2 is O, no more than one of D_3, D_4, J_3, J_4 and K_2 is S, one of D_3, D_4, J_3, J_4 and K_2 must be a covalent bond when two of D_3, D_4, J_3, J_4 and K_2 are O and S, and no more than four of D_3, D_4, J_3, J_4 and K_2 are N;

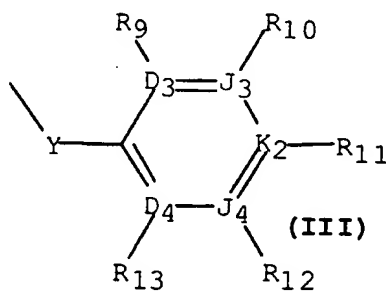
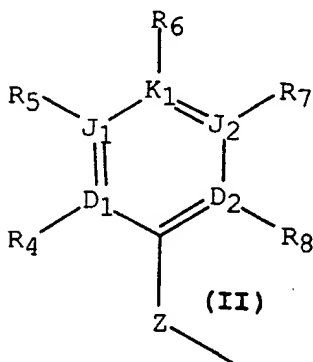
D_1, D_2, J_1, J_2 and K_1 are independently selected from the group consisting of C, O, S, N and covalent bond with the provisos that D_3, D_4, J_3, J_4 and K_2 are each carbon and at least one of D_1, D_2, J_1, J_2 and K_1 is selected from the group consisting of O, S, and N wherein, when D_1, D_2, J_1, J_2 and K_1 are selected from the group consisting of C, O, S, covalent bond, and N, no more than one of D_1, D_2, J_1, J_2 and K_1 is a covalent bond, no more than one of D_1, D_2, J_1, J_2 and K_1 is O, no more than one of D_1, D_2, J_1, J_2 and K_1 is S, one of D_1, D_2, J_1, J_2 and K_1 must be a covalent bond when two of D_1, D_2, J_1, J_2 and K_1 are O and S, and no more than four of D_1, D_2, J_1, J_2 and K_1 are N;

n is the integer 1;

X is oxy;

R_{16} is taken together with R_4 , R_8 , R_9 , or R_{13} to form a spacer selected from the group consisting of a covalent single bond, CH_2 , $CH(CH_3)$, CF_2 , $C(O)$, $C(S)$, and SO_2 ;

R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);



R_2 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, vinyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$;

R_3 is selected from the group consisting of hydrido, methyl, ethyl, vinyl, phenyl, 4-trifluoromethylphenyl, , trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and

pentafluoroethyl with the provisos that $(\text{CHR}_3)_n\text{-N(A)Q}$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 :

Y is selected from the group consisting of covalent single bond, oxy, methyleneoxy, methylene, and ethylene;

Z is selected from the group consisting of covalent single bond, oxy, methyleneoxy, methylene, and ethylene;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

R_5 and R_{10} are independently selected from the group consisting of 4-aminophenoxy, benzoyl, benzyl, benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-bromo-2-nitrophenoxy, 3-bromobenzyloxy, 4-bromobenzyloxy, 4-bromophenoxy, 5-bromopyrid-2-yloxy, 4-butoxyphenoxy, chloro, 3-chlorobenzyl, 2-chlorophenoxy, 4-chlorophenoxy, 4-chloro-3-ethylphenoxy, 3-chloro-4-fluorobenzyl, 3-chloro-4-fluorophenyl, 3-chloro-2-fluorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy, 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy, 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy, 2-cyanopyrid-3-yloxy, 4-cyanophenoxy, cyclobutoxy, cyclobutyl, cyclohexoxy, cyclohexylmethoxy, cyclopentoxy, cyclopentyl, cyclopentylcarbonyl, cyclopropyl, cyclopropylmethoxy, cyclopropoxy, 2,3-dichlorophenoxy, 2,4-dichlorophenoxy, 2,4-dichlorophenyl, 3,5-dichlorophenyl, 3,5-dichlorobenzyl, 3,4-dichlorophenoxy, 3,4-difluorophenoxy, 2,3-difluorobenzyloxy, 2,4-difluorobenzyloxy, 3,4-difluorobenzyloxy, 2,5-difluorobenzyloxy, 3,5-difluorophenoxy,

- 3,4-difluorophenyl, 3,5-difluorobenzoyloxy, 4-difluoromethoxybenzyloxy,
2,3-difluorophenoxy, 2,4-difluorophenoxy, 2,5-difluorophenoxy,
3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy, 3,5-dimethylphenoxy,
3,4-dimethylphenoxy, 3,4-dimethylbenzyl, 3,4-dimethylbenzyloxy,
5 3,5-dimethylbenzyloxy, 2,2-dimethylpropoxy, 1,3-dioxan-2-yl,
1,4-dioxan-2-yl, 1,3-dioxolan-2-yl, ethoxy, 4-ethoxyphenoxy,
4-ethylbenzyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy,
3-ethyl-5-methylphenoxy, fluoro, 4-fluoro-3-methylbenzyl,
4-fluoro-3-methylphenyl, 4-fluoro-3-methylbenzoyl, 4-fluorobenzoyloxy,
10 2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy,
3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzyloxy,
3-fluoro-5-trifluoromethylbenzyloxy, 4-fluoro-2-trifluoromethylbenzyloxy,
4-fluoro-3-trifluoromethylbenzyloxy, 2-fluorophenoxy, 4-fluorophenoxy,
2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzoyloxy,
15 4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy,
4-fluoropyrid-2-yloxy, 2-furyl, 3-furyl, heptafluoropropyl,
1,1,1,3,3,3-hexafluoropropyl, 2-hydroxy-3,3,3-trifluoropropoxy,
3-iodobenzoyloxy, isobutyl, isobutylamino, isobutoxy, 3-isoxazolyl,
4-isoxazolyl, 5-isoxazolyl, isopropoxy, isopropyl, 4-isopropylbenzyloxy,
20 3-isopropylphenoxy, 4-isopropylphenoxy, isopropylthio,
4-isopropyl-3-methylphenoxy, 3-isothiazolyl, 4-isothiazolyl,
5-isothiazolyl, 3-methoxybenzyl, 4-methoxycarbonylbutoxy,
3-methoxycarbonylprop-2-enyloxy, 4-methoxyphenyl,
3-methoxyphenylamino, 4-methoxyphenylamino, 3-methylbenzyloxy,
25 4-methylbenzyloxy, 3-methylphenoxy, 3-methyl-4-methylthiophenoxy,
4-methylphenoxy, 1-methylpropoxy, 2-methylpyrid-5-yloxy,
4-methylthiophenoxy, 2-naphthyloxy, 2-nitrophenoxy, 4-nitrophenoxy,
3-nitrophenyl, 4-nitrophenylthio, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl,
pentafluoroethyl, pentafluoroethylthio, 2,2,3,3,3-pentafluoropropyl,

- 1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl, phenoxy,
 phenylamino, 1-phenylethoxy, phenylsulfonyl, 4-propanoylphenoxy,
 propoxy, 4-propylphenoxy, 4-propoxyphenoxy, thiophen-3-yl, *sec*-butyl,
 4-*sec*-butylphenoxy, *tert*-butoxy, 3-*tert*-butylphenoxy, 4-*tert*-butylphenoxy,
 5 1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
 2-(5,6,7,8-tetrahydronaphthyl)oxy, thiazol-2-yl, thiazol-4-yl, thiazol-5-yl,
 thiophen-2-yl, 2,3,5-trifluorobenzyl, 2,2,2-trifluoroethoxy,
 2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
 3-trifluoromethoxybenzyl, 4-trifluoromethoxybenzyl,
 10 3-trifluoromethoxyphenoxy, 4-trifluoromethoxyphenoxy, trifluoromethyl,
 3-trifluoromethylbenzyl, 4-trifluoromethylbenzyl,
 2,4-bis-trifluoromethylbenzyl, 1,1-bis-trifluoromethyl-1-hydroxymethyl,
 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyl,
 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
 15 3-trifluoromethylphenyl, 3-trifluoromethylthiobenzyl,
 4-trifluoromethylthiobenzyl, 2,3,4-trifluorophenoxy,
 2,3,4-trifluorophenyl, 2,3,5-trifluorophenoxy, 3,4,5-trimethylphenoxy,
 3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy,
 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, and
 20 trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of
 chloro, fluoro, hydrido, difluoromethoxy, trifluoromethyl, trifluoromethoxy,
 pentafluoroethyl, and 1,1,2,2-tetrafluoroethoxy;

- R_7 and R_{12} are independently selected from the group consisting of
 25 hydrido, fluoro, and trifluoromethyl.

17. The compound as recited in Claim 15 or a pharmaceutically acceptable
 salt thereof, wherein:

D_1, D_2, J_1, J_2 and K_1 are each carbon with the proviso that at least one of D_3, D_4, J_3, J_4 and K_2 is selected from the group consisting of O, S, and N.

wherein D_3, D_4, J_3, J_4 and K_2 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more

5 than one of D_3, D_4, J_3, J_4 and K_2 is a covalent bond, no more than one of $D_3,$

D_4, J_3, J_4 and K_2 is O, no more than one of D_3, D_4, J_3, J_4 and K_2 is S, one of

D_3, D_4, J_3, J_4 and K_2 must be a covalent bond when two of D_3, D_4, J_3, J_4 and

K_2 are O and S, and no more than four of D_3, D_4, J_3, J_4 and K_2 are N:

D_1, D_2, J_1, J_2 and K_1 are selected from the group consisting of C, O,

10 S, N and covalent bond with the provisos that D_3, D_4, J_3, J_4 and K_2 are each

carbon and at least one of D_1, D_2, J_1, J_2 and K_1 is selected from the group

consisting of O, S, and N wherein, when D_1, D_2, J_1, J_2 and K_1 are selected

from the group consisting of C, O, S, covalent bond, and N, no more than one of D_1, D_2, J_1, J_2 and K_1 is a covalent bond, no more than one of D_1, D_2, J_1, J_2

15 and K_1 is O, no more than one of D_1, D_2, J_1, J_2 and K_1 is S, one of $D_1, D_2,$

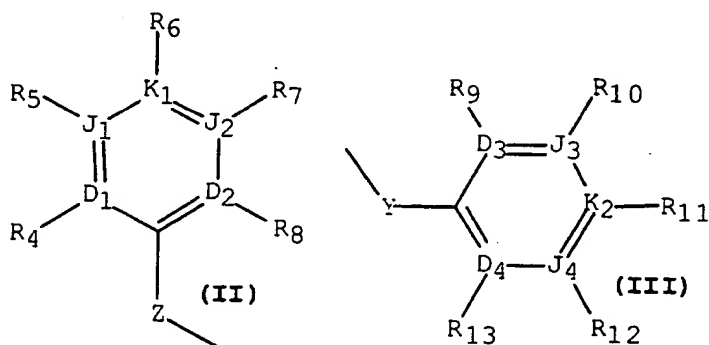
J_1, J_2 and K_1 must be a covalent bond when two of D_1, D_2, J_1, J_2 and K_1 are

O and S, and no more than four of D_1, D_2, J_1, J_2 and K_1 are N;

n is an integer selected from 1 and 2;

X is oxy;

R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III):



5

R_{16} is hydrido;

R_2 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, and haloalkoxyalkyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$:

10

R_3 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, and haloalkoxyalkyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from the group consisting of a covalent single bond, oxy and C1-C2 alkylene;

15

Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

R_4 , R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

10

18. The compound as recited in Claim 17 and pharmaceutically acceptable salts, wherein;

15

n is the integer 1;

X is oxy;

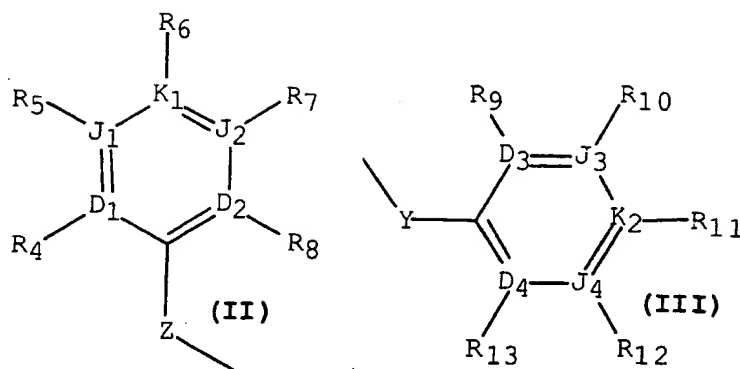
R_{16} is hydrido;

R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl,

20

chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);

264



R_2 is selected from the group consisting of hydrido, methyl, ethyl, propyl, butyl, vinyl, phenyl, 4-trifluoromethylphenyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, and

5 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$:

R_3 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, methyl, ethyl, vinyl, trifluoromethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and

10 pentafluoroethyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is selected from the group consisting of a single covalent bond, methylene, ethylene, and oxy;

15 Z is covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

- R_5 and R_{10} are independently selected from the group consisting of 4-aminophenoxy, benzoyl, benzyl, benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-bromo-2-nitrophenoxy, 3-bromobenzyloxy, 4-bromobenzyloxy, 4-bromophenoxy, 5-bromopyrid-2-yloxy, 5 4-butoxyphenoxy, chloro, 3-chlorobenzyl, 2-chlorophenoxy, 4-chlorophenoxy, 4-chloro-3-ethylphenoxy, 3-chloro-4-fluorobenzyl, 3-chloro-4-fluorophenyl, 3-chloro-2-fluorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy, 10 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy, 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy, 2-cyanopyrid-3-yloxy, 4-cyanophenoxy, cyclobutoxy, cyclobutyl, cyclohexoxy, cyclohexylmethoxy, cyclopentoxy, cyclopentyl, cyclopentylcarbonyl, cyclopropyl, cyclopropylmethoxy, cyclopropoxy, 15 2,3-dichlorophenoxy, 2,4-dichlorophenoxy, 2,4-dichlorophenyl, 3,5-dichlorophenyl, 3,5-dichlorobenzyl, 3,4-dichlorophenoxy, 3,4-difluorophenoxy, 2,3-difluorobenzyloxy, 2,4-difluorobenzyloxy, 3,4-difluorobenzyloxy, 2,5-difluorobenzyloxy, 3,5-difluorophenoxy, 3,4-difluorophenyl, 3,5-difluorobenzyloxy, 4-difluoromethoxybenzyloxy, 20 2,3-difluorophenoxy, 2,4-difluorophenoxy, 2,5-difluorophenoxy, 3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3,4-dimethylbenzyl, 3,4-dimethylbenzyloxy, 3,5-dimethylbenzyloxy, 2,2-dimethylpropoxy, 1,3-dioxan-2-yl, 1,4-dioxan-2-yl, 1,3-dioxolan-2-yl, ethoxy, 4-ethoxyphenoxy, 25 4-ethylbenzyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy, 3-ethyl-5-methylphenoxy, fluoro, 4-fluoro-3-methylbenzyl, 4-fluoro-3-methylphenyl, 4-fluoro-3-methylbenzoyl, 4-fluorobenzyloxy, 2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy, 3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzyloxy,

- 3-fluoro-5-trifluoromethylbenzyloxy. 4-fluoro-2-trifluoromethylbenzyloxy.
4-fluoro-3-trifluoromethylbenzyloxy. 2-fluorophenoxy. 4-fluorophenoxy.
2-fluoro-3-trifluoromethylphenoxy. 2-fluorobenzyloxy.
4-fluorophenylamino. 2-fluoro-4-trifluoromethylphenoxy.
5 4-fluoropyrid-2-yloxy, 2-furyl. 3-furyl. heptafluoropropyl.
1,1,1,3,3,3-hexafluoropropyl. 2-hydroxy-3,3,3-trifluoropropoxy.
3-iodobenzyloxy. isobutyl. isobutylamino. isobutoxy. 3-isoxazolyl.
4-isoxazolyl. 5-isoxazolyl. isopropoxy. isopropyl. 4-isopropylbenzyloxy,
3-isopropylphenoxy, 4-isopropylphenoxy. isopropylthio,
10 4-isopropyl-3-methylphenoxy, 3-isothiazolyl. 4-isothiazolyl.
5-isothiazolyl, 3-methoxybenzyl. 4-methoxycarbonylbutoxy.
3-methoxycarbonylprop-2-en-yloxy. 4-methoxyphenyl,
3-methoxyphenylamino, 4-methoxyphenylamino. 3-methylbenzyloxy,
4-methylbenzyloxy, 3-methylphenoxy. 3-methyl-4-methylthiophenoxy,
15 4-methylphenoxy. 1-methylpropoxy. 2-methylpyrid-5-yloxy.
4-methylthiophenoxy, 2-naphthyloxy, 2-nitrophenoxy, 4-nitrophenoxy,
3-nitrophenyl, 4-nitrophenylthio, 2-oxazolyl. 4-oxazolyl. 5-oxazolyl,
pentafluoroethyl, pentafluoroethylthio. 2,2,3,3,3-pentafluoropropyl.
1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl. phenoxy,
20 phenylamino, 1-phenylethoxy, phenylsulfonyl, 4-propanoylphenoxy,
propoxy, 4-propylphenoxy, 4-propoxyphenoxy, thiophen-3-yl. *sec*-butyl,
4-*sec*-butylphenoxy, *tert*-butoxy, 3-*tert*-butylphenoxy, 4-*tert*-butylphenoxy,
1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
2-(5,6,7,8-tetrahydronaphthyloxy), thiazol-2-yl, thiazol-4-yl. thiazol-5-yl,
25 thiophen-2-yl, 2,3,5-trifluorobenzyloxy, 2,2,2-trifluoroethoxy,
2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
3-trifluoromethoxyphenoxy. 4-trifluoromethoxyphenoxy. trifluoromethyl,
3-trifluoromethylbenzyloxy, 4-trifluoromethylbenzyloxy,
30 2,4-bis-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,

- 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
 3-trifluoromethylphenyl, 3-trifluoromethylthiobenzyloxy,
 4-trifluoromethylthiobenzyloxy, 2,3,4-trifluorophenoxy,
 5 2,3,4-trifluorophenyl, 2,3,5-trifluorophenoxy, 3,4,5-trimethylphenoxy,
 3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy,
 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, and
 trifluoromethylthio;

- R_6 and R_{11} are independently selected from the group consisting of
 10 chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy,
 trifluoromethyl, and trifluoromethoxy;

R_7 and R_{12} are independently selected from the group consisting of
 hydrido, fluoro, and trifluoromethyl.

15

19. The compound as recited in Claim 18 or a pharmaceutically acceptable
 salt thereof, wherein;

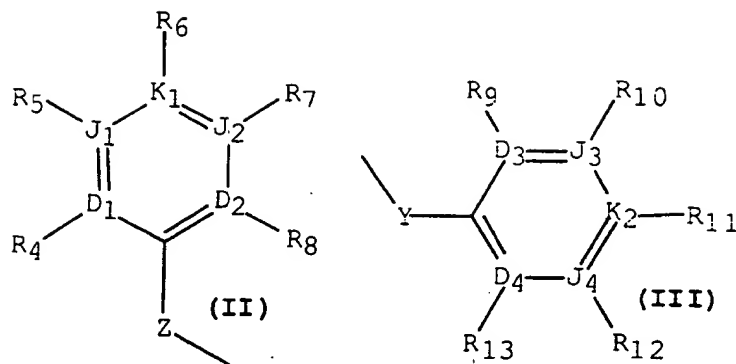
n is the integer 1;

20

X is oxy;

- R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-
 tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl,
 chlorodifluoromethyl, and pentafluoroethyl with the proviso that R_1 has a
 higher Cahn-Ingold-Prelog stereochemical system ranking than both R_2 and
 25 $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);

268



R_{16} is hydrido;

R_2 is selected from the group consisting of hydrido, methyl, ethyl, phenyl, 4-trifluoromethylphenyl, trifluoromethoxymethyl,

- 5 1,1,2,2-tetrafluoroethoxymethyl, difluoromethyl, and 2,2,3,3,3-pentafluoropropyl with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$;

R_3 is selected from the group consisting of hydrido, phenyl, 4-trifluoromethylphenyl, methyl, trifluoromethyl, difluoromethyl, and
 10 chlorodifluoromethyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 ;

Y is methylene;

Z is covalent single bond;

- 15 R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

- R_5 and R_{10} are independently selected from the group consisting of
- benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy,
 3-bromobenzyloxy, 4-bromophenoxy, 4-butoxyphenoxy,
 3-chlorobenzyloxy, 2-chlorophenoxy, 4-chloro-3-ethylphenoxy,
 5 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy,
 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy,
 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy,
 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yl, oxy,
 cyclobutoxy, cyclobutyl, cyclohexylmethoxy, cyclopentoxo, cyclopentyl,
 10 cyclopentylcarbonyl, cyclopropylmethoxy, 2,3-dichlorophenoxy,
 2,4-dichlorophenoxy, 2,4-dichlorophenyl, 3,5-dichlorophenyl,
 3,5-dichlorobenzyl, 3,4-dichlorophenoxy, 3,4-difluorophenoxy,
 2,3-difluorobenzyloxy, 3,5-difluorobenzyloxy, difluoromethoxy,
 3,5-difluorophenoxy, 3,4-difluorophenyl, 2,3-difluorophenoxy,
 15 2,4-difluorophenoxy, 2,5-difluorophenoxy, 3,5-dimethoxyphenoxy,
 3-dimethylaminophenoxy, 3,4-dimethylbenzyloxy, 3,5-dimethylbenzyloxy,
 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 1,3-dioxolan-2-yl,
 3-ethylbenzyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy,
 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylbenzyl, 4-fluorobenzyloxy,
 20 2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy, 3-fluorophenoxy,
 3-fluoro-2-nitrophenoxy, 2-fluoro-3-trifluoromethylbenzyloxy,
 3-fluoro-5-trifluoromethylbenzyloxy, 2-fluorophenoxy, 4-fluorophenoxy,
 2-fluoro-3-trifluoromethylphenoxy, 2-fluorobenzyloxy,
 4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy, 2-furyl, 3-furyl,
 25 heptafluoropropyl, 1,1,1,3,3,3-hexafluoropropyl,
 2-hydroxy-3,3,3-trifluoropropoxy, isobutoxy, isobutyl, 3-isoxazolyl,
 4-isoxazolyl, 5-isoxazolyl, isopropoxy, 3-isopropylbenzyloxy,
 3-isopropylphenoxy, isopropylthio, 4-isopropyl-3-methylphenoxy,
 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl, 3-methoxybenzyl,

- 4-methoxyphenylamino, 3-methylbenzyloxy, 4-methylbenzyloxy,
 3-methylphenoxy, 3-methyl-4-methylthiophenoxy, 4-methylphenoxy,
 1-methylpropoxy, 2-methylpyrid-5-yloxy, 4-methylthiophenoxy,
 2-naphthylloxy, 2-nitrophenoxy, 4-nitrophenoxy, 3-nitrophenyl, 2-oxazolyl,
 5 4-oxazolyl, 5-oxazolyl, pentafluoroethyl, pentafluoroethylthio, 2,2,3,3,3-
 pentafluoropropyl, 1,1,3,3,3-pentafluoropropyl, 1,1,2,2,3-pentafluoropropyl,
 phenoxy, phenylamino, 1-phenylethoxy, 4-propylphenoxy,
 4-propoxyphenoxy, thiophen-3-yl, tert-butoxy, 3-tert-butylphenoxy,
 4-tert-butylphenoxy, 1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
 10 2-(5,6,7,8-tetrahydronaphthylloxy), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl,
 thiophen-2-yl, 2,2,2-trifluoroethoxy, 2,2,2-trifluoroethyl,
 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
 3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
 4-trifluoromethoxyphenoxy, 3-trifluoromethoxyphenoxy, trifluoromethyl,
 15 3-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,
 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
 3-trifluoromethylphenyl, 2,3,4-trifluorophenoxy, 2,3,5-trifluorophenoxy,
 3,4,5-trimethylphenoxy, 3-difluoromethoxyphenoxy,
 20 3-pentafluoroethylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy,
 3-trifluoromethylthiophenoxy, 3-trifluoromethylthiobenzyloxy, and
 trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of
 chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy, and
 25 trifluoromethyl;

R_7 and R_{12} are independently selected from the group consisting of
 hydrido, fluoro, and trifluoromethyl.

20. The compound as recited in Claim 17 or a pharmaceutically acceptable salt, wherein:

D_1, D_2, J_1, J_2 and K_1 are each carbon:

D_3, D_4, J_3, J_4 and K_2 are independently selected from the group

- 5 consisting of C, N, O, S and covalent bond with the provisos that no more than one of D_3, D_4, J_3, J_4 and K_2 is a covalent bond, no more than one of D_3, D_4, J_3, J_4 and K_2 is O, no more than one of D_3, D_4, J_3, J_4 and K_2 is S, one of D_3, D_4, J_3, J_4 and K_2 must be a covalent bond when two of D_3, D_4, J_3, J_4 and K_2 are O and S, no more than four of D_3, D_4, J_3, J_4 and K_2 are N, and one of
- 10 D_3, D_4, J_3, J_4 and K_2 is selected from the group consisting of O, S, and N:

n is the integer 1;

X is oxy;

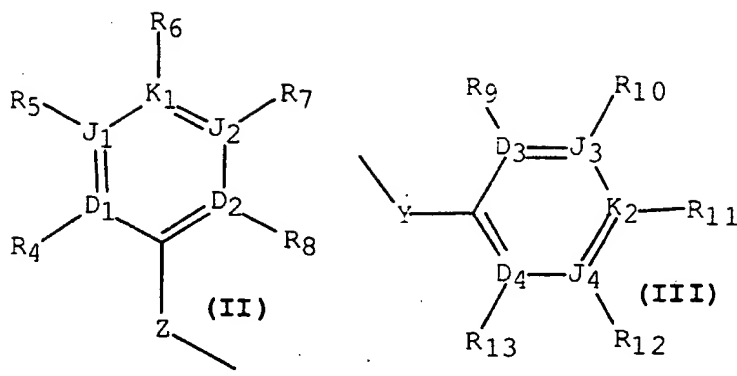
R_{16} is hydrido;

- 15 R_1 is haloalkyl with the proviso that R_1 has a higher Cahn-Ingold-

Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$

wherein A is Formula (II) and Q is Formula (III);

272



R_2 is selected from the group consisting of hydrido, alkyl, aryl,

haloalkyl, and haloalkoxy with the proviso that R_2 has a lower Cahn-Ingold-

Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$:

- 5 R_3 is selected from the group consisting of hydrido, alkyl, and haloalkyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 :

Y is a C1-C2 alkylene;

- 10 Z is covalent single bond;

R_{14} is hydrido;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

- 15 R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy,

cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, and heteroaryloxyalkyl.

- 5 21. The compound as recited in Claim 17 or a pharmaceutically acceptable salt, wherein:

D_3 , D_4 , J_3 , J_4 and K_2 are each carbon;

D_1 , D_2 , J_1 , J_2 and K_1 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that no more

- 10 than one of D_1 , D_2 , J_1 , J_2 and K_1 is a covalent bond, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 is O, no more than one of D_1 , D_2 , J_1 , J_2 and K_1 is S, one of D_1 , D_2 , J_1 , J_2 and K_1 must be a covalent bond when two of D_1 , D_2 , J_1 , J_2 and K_1 are O and S, no more than four of D_1 , D_2 , J_1 , J_2 and K_1 are N, and one of D_1 , D_2 , J_1 , J_2 and K_1 is selected from the group consisting of O, S, and N;

15

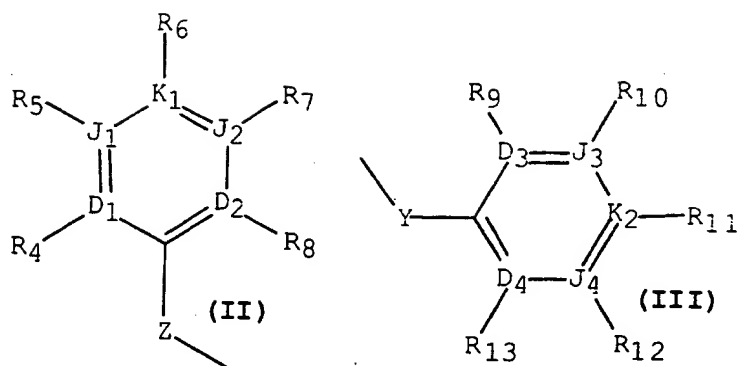
n is the integer 1;

X is oxy;

R_{16} is hydrido;

R_1 is haloalkyl with the proviso that R_1 has a higher Cahn-Ingold-

- 20 Prelog stereochemical system ranking than both R_2 and $(CHR_3)_n-N(A)Q$ wherein A is Formula (II) and Q is Formula (III);



R_2 is selected from the group consisting of hydrido, alkyl, aryl,

haloalkyl, and haloalkoxy with the proviso that R_2 has a lower Cahn-Ingold-Prelog system ranking than both R_1 and $(CHR_3)_n-N(A)Q$:

- 5 R_3 is selected from the group consisting of hydrido, alkyl, and haloalkyl with the provisos that $(CHR_3)_n-N(A)Q$ has a lower Cahn-Ingold-Prelog stereochemical system ranking than R_1 and a higher Cahn-Ingold-Prelog stereochemical system ranking than R_2 :

Y is a C1-C2 alkylene:

- 10 Z is covalent single bond:

R_{14} is hydrido:

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

- 15 R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy.

275

cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, and heteroaryloxyalkyl.

22. The compound as recited in any one of Claims 20 or 21 or a
 5 pharmaceutically acceptable salt thereof, wherein;

n is the integer 1;

X is oxy;

- R_1 is selected from the group consisting of trifluoromethyl and
 10 pentafluoroethyl;

R_{16} is hydrido;

R_2 is hydrido;

R_3 is selected from the group consisting of hydrido,
 methyl, trifluoromethyl, and difluoromethyl;

- 15 Y is methylene;
 Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group
 consisting of hydrido and fluoro;

- R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy,
 20 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-
 difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy,
 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy,
 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 3-
 pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-
 25 tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyloxy),
 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy,

276

3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy:

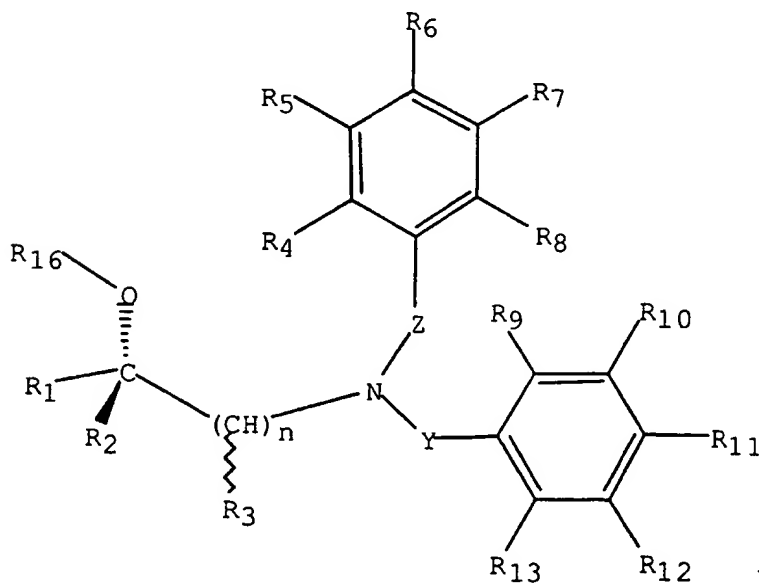
R_{10} is selected from the group consisting of cyclopentyl, 1,1,2,2-tetrafluoroethoxy, 2-furyl, 1,1-bis-trifluoromethyl-1-hydroxymethyl, isobutyl, isopropoxy, pentafluoroethyl, trifluoromethoxy, trifluoromethyl, and trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of fluoro and hydrido;

R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

10

23. A compound having the formula:



or a pharmaceutically acceptable salt thereof, wherein:

15

n is an integer selected from 1 and 2;

R_1 is selected from the group consisting of haloalkyl and haloalkoxyalkyl;

R_{16} is hydrido;

R_2 is hydrido;

5 R_3 is hydrido;

Y is selected from the group consisting of a covalent single bond and C1-C2 alkylene;

Z is selected from the group consisting of a covalent single bond and C1-C2 alkylene;

10

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the
 15 group consisting of perhaloaryloxy, N-aryl-N-alkylamino, heterocyclalkoxy, heterocyclthio, hydroxyalkoxy, carboxamidoalkoxy, alkoxy-carbonylalkoxy, alkoxy-carbonylalkenyloxy, aralkanoylalkoxy, aralkenoyl, N-arylcarboxamidoalkoxy, cycloalkylcarbonyl, cyanoalkoxy, heterocyclcarbonyl, hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl,
 20 alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, heterocyclloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

25

24. The compound as recited in Claim 23 or a pharmaceutically acceptable salt thereof wherein:

5 n is the integer 1;

R_{16} is hydrido;

R_1 is selected from the group consisting of trifluoromethyl, 1,1,2,2-tetrafluoroethoxymethyl, trifluoromethoxymethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl;

10 R_2 is hydrido;

R_3 is hydrido;

 Y is selected from the group consisting of methylene, and ethylene;

 Z is selected from the group consisting of covalent single bond and methylene;

15 R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro;

R_5 and R_{10} are independently selected from the group consisting of 4-aminophenoxy, benzoyl, benzyl, benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 4-bromo-2-nitrophenoxy, 3-bromobenzyloxy, 20 4-bromobenzyloxy, 4-bromophenoxy, 5-bromopyrid-2-yloxy, 4-butoxyphenoxy, chloro, 3-chlorobenzyl, 2-chlorophenoxy, 4-chlorophenoxy, 4-chloro-3-ethylphenoxy, 3-chloro-4-fluorobenzyl, 3-chloro-4-fluorophenyl, 3-chloro-2-fluorobenzyloxy, 3-chlorobenzyloxy, 4-chlorobenzyloxy, 4-chloro-3-methylphenoxy, 2-chloro-4-fluorophenoxy, 25 4-chloro-2-fluorophenoxy, 4-chlorophenoxy, 3-chloro-4-ethylphenoxy, 3-chloro-4-methylphenoxy, 3-chloro-4-fluorophenoxy,

- 4-chloro-3-fluorophenoxy, 4-chlorophenylamino, 5-chloropyrid-3-yloxy,
2-cyanopyrid-3-yloxy, 4-cyanophenoxy, cyclobutoxy, cyclobutyl,
cyclohexoxy, cyclohexylmethoxy, cyclopentoxy, cyclopentyl,
cyclopentylcarbonyl, cyclopropyl, cyclopropylmethoxy, cyclopropoxy.
- 5 2,3-dichlorophenoxy, 2,4-dichlorophenoxy, 2,4-dichlorophenyl,
3,5-dichlorophenyl, 3,5-dichlorobenzyl, 3,4-dichlorophenoxy,
3,4-difluorophenoxy, 2,3-difluorobenzoyloxy, 2,4-difluorobenzoyloxy,
3,4-difluorobenzoyloxy, 2,5-difluorobenzoyloxy, 3,5-difluorophenoxy,
3,4-difluorophenyl, 3,5-difluorobenzoyloxy, 4-difluoromethoxybenzoyloxy,
- 10 2,3-difluorophenoxy, 2,4-difluorophenoxy, 2,5-difluorophenoxy,
3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy, 3,5-dimethylphenoxy,
3,4-dimethylphenoxy, 3,4-dimethylbenzyl, 3,4-dimethylbenzoyloxy,
3,5-dimethylbenzoyloxy, 2,2-dimethylpropoxy, 1,3-dioxan-2-yl,
1,4-dioxan-2-yl, 1,3-dioxolan-2-yl, ethoxy, 4-ethoxyphenoxy,
- 15 4-ethylbenzoyloxy, 3-ethylphenoxy, 4-ethylaminophenoxy,
3-ethyl-5-methylphenoxy, fluoro, 4-fluoro-3-methylbenzyl,
4-fluoro-3-methylphenyl, 4-fluoro-3-methylbenzoyl, 4-fluorobenzoyloxy,
2-fluoro-3-methylphenoxy, 3-fluoro-4-methylphenoxy,
3-fluorophenoxy, 3-fluoro-2-nitrophenoxy,
- 20 2-fluoro-3-trifluoromethylbenzoyloxy, 3-fluoro-5-trifluoromethylbenzoyloxy,
4-fluoro-2-trifluoromethylbenzoyloxy, 4-fluoro-3-trifluoromethylbenzoyloxy,
2-fluorophenoxy, 4-fluorophenoxy, 2-fluoro-3-trifluoromethylphenoxy,
2-fluorobenzoyloxy, 4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy,
4-fluoropyrid-2-yloxy, 2-furyl, 3-furyl, heptafluoropropyl, 1,1,1,3,3,3-
- 25 hexafluoropropyl, 2-hydroxy-3,3,3-trifluoropropoxy, 3-iodobenzoyloxy,
isobutyl, isobutylamino, isobutoxy, 3-isoxazolyl, 4-isoxazolyl, 5-isoxazolyl,
isopropoxy, isopropyl, 4-isopropylbenzoyloxy, 3-isopropylphenoxy,
4-isopropylphenoxy, isopropylthio, 4-isopropyl-3-methylphenoxy,
3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl, 3-methoxybenzyl,
- 30 4-methoxycarbonylbutoxy, 3-methoxycarbonylprop-2-enyloxy,

- 4-methoxyphenyl, 3-methoxyphenylamino, 4-methoxyphenylamino,
3-methylbenzyloxy, 4-methylbenzyloxy, 3-methylphenoxy,
3-methyl-4-methylthiophenoxy, 4-methylphenoxy, 1-methylpropoxy,
2-methylpyrid-5-yloxy, 4-methylthiophenoxy, 2-naphthyloxy,
5 2-nitrophenoxy, 4-nitrophenoxy, 3-nitrophenyl, 4-nitrophenylthio, 2-
oxazolyl, 4-oxazolyl, 5-oxazolyl, pentafluoroethyl, pentafluoroethylthio,
2,2,3,3,3-pentafluoropropyl, 1,1,3,3,3-pentafluoropropyl,
1,1,2,2,3-pentafluoropropyl, phenoxy, phenylamino, 1-phenylethoxy,
phenylsulfonyl, 4-propanoylphenoxy, propoxy, 4-propylphenoxy,
10 4-propoxyphenoxy, thiophen-3-yl, *sec*-butyl, 4-*sec*-butylphenoxy,
tert-butoxy, 3-*tert*-butylphenoxy, 4-*tert*-butylphenoxy,
1,1,2,2-tetrafluoroethoxy, tetrahydrofuran-2-yl,
2-(5,6,7,8-tetrahydronaphthyloxy), thiazol-2-yl, thiazol-4-yl, thiazol-5-yl,
thiophen-2-yl, 2,3,5-trifluorobenzyloxy, 2,2,2-trifluoroethoxy,
15 2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
3-trifluoromethoxyphenoxy, 4-trifluoromethoxyphenoxy, trifluoromethyl, 3-
trifluoromethylbenzyloxy, 4-trifluoromethylbenzyloxy,
2,4-bis-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,
20 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
3-trifluoromethylphenyl, 3-trifluoromethylthiobenzyloxy,
4-trifluoromethylthiobenzyloxy, 2,3,4-trifluorophenoxy,
2,3,4-trifluorophenyl, 2,3,5-trifluorophenoxy, 3,4,5-trimethylphenoxy,
25 3-difluoromethoxyphenoxy, 3-pentafluoroethylphenoxy,
3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 3-trifluoromethylthiophenoxy, and
trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy, trifluoromethyl, and trifluoromethoxy;

R_7 and R_{12} are independently selected from the group consisting of
5 hydrido, fluoro, and trifluoromethyl.

25. The compound as recited in Claim 24 or a pharmaceutically acceptable salt thereof, wherein;

10

n is the integer 1;

R_1 is selected from the group consisting of trifluoromethyl, difluoromethyl, chlorodifluoromethyl, and pentafluoroethyl;

R_{16} is hydrido;

15

R_2 is hydrido;

R_3 is hydrido;

Y is methylene;

Z is covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group
20 consisting of hydrido and fluoro;

R_5 and R_{10} are independently selected from the group consisting of benzyloxy, 5-bromo-2-fluorophenoxy, 4-bromo-3-fluorophenoxy, 3-bromobenzyloxy, 4-bromophenoxy, 4-butoxyphenoxy, 3-chlorobenzyloxy, 2-chlorophenoxy, 4-chloro-3-ethylphenoxy, 4-chloro-3-methylphenoxy,
25 2-chloro-4-fluorophenoxy, 4-chloro-2-fluorophenoxy, 4-chlorophenoxy,

- 3-chloro-4-ethylphenoxy, 3-chloro-4-methylphenoxy,
 3-chloro-4-fluorophenoxy, 4-chloro-3-fluorophenoxy,
 4-chlorophenylamino, 5-chloropyrid-3-yloxy, cyclobutoxy, cyclobutyl,
 cyclohexylmethoxy, cyclopentoxo, cyclopentyl, cyclopentylcarbonyl,
 5 cyclopropylmethoxy, 2,3-dichlorophenoxy, 2,4-dichlorophenoxy,
 2,4-dichlorophenyl, 3,5-dichlorophenyl, 3,5-dichlorobenzyl,
 3,4-dichlorophenoxy, 3,4-difluorophenoxy, 2,3-difluorobenzoyloxy,
 3,5-difluorobenzoyloxy, difluoromethoxy, 3,5-difluorophenoxy,
 3,4-difluorophenyl, 2,3-difluorophenoxy, 2,4-difluorophenoxy,
 10 2,5-difluorophenoxy, 3,5-dimethoxyphenoxy, 3-dimethylaminophenoxy,
 3,4-dimethylbenzoyloxy, 3,5-dimethylbenzoyloxy, 3,5-dimethylphenoxy,
 3,4-dimethylphenoxy, 1,3-dioxolan-2-yl, 3-ethylbenzoyloxy,
 3-ethylphenoxy, 4-ethylaminophenoxy, 3-ethyl-5-methylphenoxy,
 4-fluoro-3-methylbenzyl, 4-fluorobenzoyloxy, 2-fluoro-3-methylphenoxy,
 15 3-fluoro-4-methylphenoxy, 3-fluorophenoxy, 3-fluoro-2-nitrophenoxy,
 2-fluoro-3-trifluoromethylbenzoyloxy, 3-fluoro-5-trifluoromethylbenzoyloxy,
 2-fluorophenoxy, 4-fluorophenoxy, 2-fluoro-3-trifluoromethylphenoxy,
 2-fluorobenzoyloxy, 4-fluorophenylamino, 2-fluoro-4-trifluoromethylphenoxy,
 2-furyl, 3-furyl, heptafluoropropyl, 1,1,1,3,3,3-hexafluoropropyl,
 20 2-hydroxy-3,3,3-trifluoropropoxy, isobutoxy, isobutyl, 3-isoxazolyl,
 4-isoxazolyl, 5-isoxazolyl, isopropoxy, 3-isopropylbenzoyloxy,
 3-isopropylphenoxy, isopropylthio, 4-isopropyl-3-methylphenoxy,
 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl, 3-methoxybenzyl,
 4-methoxyphenylamino, 3-methylbenzoyloxy, 4-methylbenzoyloxy,
 25 3-methylphenoxy, 3-methyl-4-methylthiophenoxy, 4-methylphenoxy,
 1-methylpropoxy, 2-methylpyrid-5-yloxy, 4-methylthiophenoxy,
 2-naphthylloxy, 2-nitrophenoxy, 4-nitrophenoxy, 3-nitrophenyl, 2-oxazolyl,
 4-oxazolyl, 5-oxazolyl, pentafluoroethyl, pentafluoroethylthio,
 2,2,3,3,3-pentafluoropropyl, 1,1,1,3,3,3-pentafluoropropyl,
 30 1,1,2,2,3-pentafluoropropyl, phenoxy, phenylamino, 1-phenylethoxy,

- 4-propylphenoxy, 4-propoxyphenoxy, thiophen-3-yl,tert -butoxy,
 3-tert -butylphenoxy, 4-tert -butylphenoxy, 1,1,2,2-tetrafluoroethoxy,
 tetrahydrofuran-2-yl, 2-(5,6,7,8-tetrahydronaphthyl)oxy, thiazol-2-yl,
 thiazol-4-yl, thiazol-5-yl, thiophen-2-yl, 2,2,2-trifluoroethoxy,
 5 2,2,2-trifluoroethyl, 3,3,3-trifluoro-2-hydroxypropyl, trifluoromethoxy,
 3-trifluoromethoxybenzyloxy, 4-trifluoromethoxybenzyloxy,
 4-trifluoromethoxyphenoxy, 3-trifluoromethoxyphenoxy, trifluoromethyl,
 3-trifluoromethylbenzyloxy, 1,1-bis-trifluoromethyl-1-hydroxymethyl,
 3-trifluoromethylbenzyl, 3,5-bis-trifluoromethylbenzyloxy,
 10 4-trifluoromethylphenoxy, 3-trifluoromethylphenoxy,
 3-trifluoromethylphenyl, 2,3,4-trifluorophenoxy, 2,3,5-trifluorophenoxy,
 3,4,5-trimethylphenoxy, 3-difluoromethoxyphenoxy,
 3-pentafluoroethylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy,
 3-trifluoromethylthiophenoxy, 3-trifluoromethylthiobenzyloxy, and
 15 trifluoromethylthio;

R_6 and R_{11} are independently selected from the group consisting of
 chloro, fluoro, hydrido, pentafluoroethyl, 1,1,2,2-tetrafluoroethoxy, and
 trifluoromethyl;

- R_7 and R_{12} are independently selected from the group consisting of
 20 hydrido, fluoro, and trifluoromethyl.

26. The compound as recited in Claim 23 or a pharmaceutically acceptable
 salt, wherein;

- 25 n is the integer 1;

R_{16} is hydrido;

R_1 is haloalkyl;

R_2 is is hydrido;

R_3 is is hydrido;

Y is methylene;

Z is a covalent single bond;

5 R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of perhaloaryloxy, N-aryl-N-alkylamino, heterocyclylalkoxy, heterocyclylthio, hydroxyalkoxy, aralkanoylalkoxy, aralkenoyl, cycloalkylcarbonyl, cyanoalkoxy, heterocyclylcarbonyl, hydrido, 10 alkyl, halo, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, and heteroaryloxyalkyl.

15

27. The compound as recited in Claim 26 or a pharmaceutically acceptable salt, wherein;

n is the integer 1;

20

R_{16} is hydrido;

R_1 is haloalkyl;

R_2 is is hydrido;

R_3 is is hydrido;

Y is methylene;

Z is a covalent single bond:

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and halo:

R_5 is selected from the group consisting of perhaloaryloxy.

- 5 N-aryl-N-alkylamino, heterocyclalkoxy, heterocyclthio, hydroxyalkoxy, aralkanoylalkoxy, aralkenoyl, cycloalkylcarbonyl, cyanoalkoxy, heterocyclcarbonyl, haloalkyl, haloalkoxy, aryl, alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy, heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy, cycloalkylalkanoyl, heteroaryl,
- 10 cycloalkyl, haloalkylthio, hydroxyhaloalkyl, heteroaralkoxy, and heteroaryloxyalkyl.

R_{10} is selected from the group consisting of haloalkyl, haloalkoxy, aryl, alkylthio, alkoxy, aralkyl, alkyl, cycloalkoxy, cycloalkylalkoxy, heteroaryl, cycloalkyl, haloalkylthio, and hydroxyhaloalkyl.

- 15 R_6 and R_{11} are independently selected from the group consisting of hydrido and halo;

R_7 and R_{12} are independently selected from the group consisting of hydrido and halo.

- 20 28. The compound as recited in Claim 27 or a pharmaceutically acceptable salt thereof, wherein;

n is the integer 1;

R_1 is trifluoromethyl;

- 25 R_{16} is hydrido;

R_2 is hydrido:

R_3 is hydrido:

Y is methylene:

Z is a covalent single bond:

- 5 R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro:

- R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy, 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 10 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy, 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyloxy), 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 15 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R_{10} is selected from the group consisting of cyclopentyl, 1,1,2,2-tetrafluoroethoxy, 2-furyl, 1,1-bis-trifluoromethyl-1-hydroxymethyl, pentafluoroethyl, trifluoromethoxy, trifluoromethyl, and trifluoromethylthio;

- R_6 and R_{11} are independently selected from the group consisting of 20 fluoro and hydrido:

R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

29. The compound as recited in Claim 28 or a pharmaceutically 25 acceptable salt thereof, wherein:

n is the integer 1:

R_1 is trifluoromethyl:

R_{16} is hydrido:

R_2 is hydrido:

5 R_3 is hydrido:

Y is methylene:

Z is a covalent single bond:

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido and fluoro:

10 R_5 is selected from the group consisting of 5-bromo-2-fluorophenoxy, 4-chloro-3-ethylphenoxy, 2,3-dichlorophenoxy, 3,4-dichlorophenoxy, 3-difluoromethoxyphenoxy, 3,5-dimethylphenoxy, 3,4-dimethylphenoxy, 3-ethylphenoxy, 3-ethyl-5-methylphenoxy, 4-fluoro-3-methylphenoxy, 4-fluorophenoxy, 3-isopropylphenoxy, 3-methylphenoxy, 15 3-pentafluoroethylphenoxy, 3-tert-butylphenoxy, 3-(1,1,2,2-tetrafluoroethoxy)phenoxy, 2-(5,6,7,8-tetrahydronaphthyl)oxy, 3-trifluoromethoxybenzyloxy, 3-trifluoromethoxyphenoxy, 3-trifluoromethylbenzyloxy, and 3-trifluoromethylthiophenoxy;

R_{10} is selected from the group consisting of 1,1,2,2-tetrafluoroethoxy, 20 pentafluoroethyl, and trifluoromethyl;

R_6 and R_{11} are independently selected from the group consisting of fluoro and hydrido;

R_7 and R_{12} are independently selected from the group consisting of hydrido and fluoro.

30. A compound as recited in Claim 23 or a pharmaceutically acceptable salt thereof wherein said compound is selected from the group consisting of:

- 5 (2R)-3-[[3-(3-trifluoromethoxyphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-isopropylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-cyclopropylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 10 (2R)-3-[[3-(3-(2-furyl)phenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2,3-dichlorophenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 15 (2R)-3-[[3-(4-fluorophenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-methylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2-fluoro-5-bromophenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 20 (2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-[3-(1,1,2,2-tetrafluoroethoxy)phenoxy]phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 25 (2R)-3-[[3-[3-(pentafluoroethyl)phenoxy]phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3,5-dimethylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-ethylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 30 phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

- (2R)-3-[[3-(3-*i*-butylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(3-methylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- 5 (2R)-3-[[3-(5,6,7,8-tetrahydro-2-naphthoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(phenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(*N,N*-dimethylamino)phenoxy]phenyl][[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- 10 (2R)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl][3-[[3-(trifluoromethoxy)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl][3-[[3-(trifluoromethyl)phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:
- 15 (2R)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl][3-[[3,5-dimethylphenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl][3-[[3-(trifluoromethylthio)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl][3-[[3,5-difluorophenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:
- 20 3-[[[3-(1,1,2,2-tetrafluoroethoxy)phenyl]-methyl][3-(cyclohexylmethoxy)-phenyl]amino]-1,1,1-trifluoro-2-propanol:
- 3-[[3-(2-difluoromethoxy-4-pyridyloxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- 25 (2R)-3-[[3-(2-trifluoromethyl-4-pyridyloxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(3-difluoromethoxyphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[[3-(3-trifluoromethylthio)phenoxy]phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- 30

(2R)-3-[[3-(4-chloro-3-trifluoromethylphenoxy)phenyl][[3-(1,1,2,2-tetrafluoroethoxy)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(3-trifluoromethoxyphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

5 (2R)-3-[[3-(3-isopropylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(3-cyclopropylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

10 (2R)-3-[[3-(3-(2-furyl)phenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(2,3-dichlorophenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(4-fluorophenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

15 (2R)-3-[[3-(4-methylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(2-fluoro-5-bromophenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

20 (2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-[3-(1,1,2,2-tetrafluoroethoxy)phenoxy]phenyl][[3-(pentafluoroethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-[3-(pentafluoroethyl)phenoxy]phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

25 (2R)-3-[[3-(3,5-dimethylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(3-ethylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

30 (2R)-3-[[3-(3-*t*-butylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

- (2R)-3-[[3-(3-methylphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(5,6,7,8-tetrahydro-2-naphthoxy)phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 5 (2R)-3-[[3-(phenoxy)phenyl][[3-(pentafluoroethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-[3-(*N,N*-dimethylamino)phenoxy]phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[3-(pentafluoroethyl)phenyl]methyl][3-[[3-(trifluoromethoxy)phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- 10 (2R)-3-[[[3-(pentafluoroethyl)phenyl]methyl][3-[[3-(trifluoromethyl)-phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[3-(pentafluoroethyl)phenyl]methyl][3-[[3,5-dimethylphenyl]methoxy]-phenyl]amino]-1,1,1-trifluoro-2-propanol;
- 15 (2R)-3-[[[3-(pentafluoroethyl)phenyl]methyl][3-[[3-(trifluoromethylthio)phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[3-(pentafluoroethyl)phenyl]methyl][3-[[3,5-difluorophenyl]methoxy]-phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[3-(pentafluoroethyl)phenyl]methyl][3-[cyclohexylmethoxy]phenyl]-amino]-1,1,1-trifluoro-2-propanol;
- 20 (2R)-3-[[3-(2-difluoromethoxy-4-pyridyloxy)phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2-trifluoromethyl-4-pyridyloxy)phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 25 (2R)-3-[[3-(3-difluoromethoxyphenoxy)phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[3-(3-trifluoromethylthio)phenoxy]phenyl][[3-(pentafluoroethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-chloro-3-trifluoromethylphenoxy)phenyl][[3-(pentafluoroethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 30

- (2R)-3-[[3-(3-trifluoromethoxyphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-isopropylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- 5 (2R)-3-[[3-(3-cyclopropylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-(2-furyl)phenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2,3-dichlorophenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- 10 (2R)-3-[[3-(4-fluorophenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-methylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 15 (2R)-3-[[3-(2-fluoro-5-bromophenoxy)phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-[3-(1,1,2,2-tetrafluoroethoxy)phenoxy]phenyl][[3-(heptafluoropropyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 20 (2R)-3-[[3-[3-(pentafluoroethyl)phenoxy]phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3,5-dimethylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- 25 (2R)-3-[[3-(3-ethylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-*t*-butylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-methylphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 30 (heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(5,6,7,8-tetrahydro-2-naphthoxy)phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(phenoxy)phenyl][[3-(heptafluoropropyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

5 (2R)-3-[[3-[3-(*N,N*-dimethylamino)phenoxy]phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[[3-(heptafluoropropyl)phenyl]methyl][3-[[3-(trifluoromethoxy)phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;

10 (2R)-3-[[[3-(heptafluoropropyl)phenyl]methyl][3-[[3-(trifluoromethyl)phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[[3-(heptafluoropropyl)phenyl]methyl][3-[[3,5-dimethylphenyl]methoxy]-phenyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[[3-(heptafluoropropyl)phenyl]methyl][3-[[3-(trifluoromethylthio)phenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;

15 (2R)-3-[[[3-(heptafluoropropyl)phenyl]methyl][3-[[3,5-difluorophenyl]methoxy]-phenyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[[3-(heptafluoropropyl)phenyl]methyl][3-[cyclohexylmethoxy]phenyl]-amino]-1,1,1-trifluoro-2-propanol;

20 (2R)-3-[[3-(2-difluoromethoxy-4-pyridyloxy)phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(2-trifluoromethyl-4-pyridyloxy)phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(3-difluoromethoxyphenoxy)phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

25 (2R)-3-[[[3-(3-trifluoromethylthio)phenoxy]phenyl][[3-(heptafluoropropyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-(4-chloro-3-trifluoromethylphenoxy)phenyl][[3-(heptafluoropropyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

30 (2R)-3-[[3-(3-trifluoromethoxyphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

- (2R)-3-[[3-(3-isopropylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(3-cyclopropylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- 5 (2R)-3-[[3-(3-(2-furyl)phenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(2,3-dichlorophenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(4-fluorophenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- 10 (2R)-3-[[3-(4-methylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(2-fluoro-5-bromophenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:
- 15 (2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-[3-(1,1,2,2-tetrafluoroethoxy)phenoxy]phenyl][[2-fluoro-5-(trifluoro-methyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-[3-(pentafluoroethyl)phenoxy]phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:
- 20 (2R)-3-[[3-(3,5-dimethylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(3-ethylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol:
- 25 (2R)-3-[[3-(3-*t*-butylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(3-methylphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol:
- (2R)-3-[[3-(5,6,7,8-tetrahydro-2-naphthoxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:
- 30

- (2R)-3-[[3-(phenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-[3-(*N,N*-dimethylamino)phenoxy]phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 5 (2R)-3-[[[2-fluoro-5-(trifluoromethyl)phenyl]methyl][3-[[3-(trifluoromethoxy)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[2-fluoro-5-(trifluoromethyl)phenyl]methyl][3-[[3-(trifluoromethyl)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[2-fluoro-5-(trifluoromethyl)phenyl]methyl][3-[[3,5-dimethylphenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- 10 (2R)-3-[[[2-fluoro-5-(trifluoromethyl)phenyl]methyl][3-[[3-(trifluoromethylthio)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[2-fluoro-5-(trifluoromethyl)phenyl]methyl][3-[[3,5-difluorophenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol;
- 15 (2R)-3-[[[2-fluoro-5-(trifluoromethyl)phenyl]methyl][3-[cyclohexylmethoxy]-phenyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2-difluoromethoxy-4-pyridyloxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2-trifluoromethyl-4-pyridyloxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 20 (2R)-3-[[3-(3-difluoromethoxyphenoxy)phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[[3-(3-trifluoromethylthio)phenoxy]phenyl][[2-fluoro-5-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 25 (2R)-3-[[3-(4-chloro-3-trifluoromethylphenoxy)phenyl][[2-fluoro-5-(trifluoro-methyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-trifluoromethoxyphenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-isopropylphenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 30

- (2R)-3-[[3-(3-cyclopropylphenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-(2-furyl)phenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- 5 (2R)-3-[[3-(2,3-dichlorophenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-fluorophenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-methylphenoxy)phenyl][[2-fluoro-4-
- 10 (trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(2-fluoro-5-bromophenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(4-chloro-3-ethylphenoxy)phenyl][[2-fluoro-4-
- (trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- 15 (2R)-3-[[3-[3-(1,1,2,2-tetrafluoroethoxy)phenoxy]phenyl][[2-fluoro-4-(trifluoro-methyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-[3-(pentafluoroethyl)phenoxy]phenyl][[2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3,5-dimethylphenoxy)phenyl][[2-fluoro-4-
- 20 (trifluoromethyl)phenyl]-methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-ethylphenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(3-*t*-butylphenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- 25 (2R)-3-[[3-(3-methylphenoxy)phenyl][[2-fluoro-4-(trifluoromethyl)phenyl]methyl]-amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(5,6,7,8-tetrahydro-2-naphthoxy)phenyl][[2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;
- (2R)-3-[[3-(phenoxy)phenyl][[2-fluoro-4-
- 30 (trifluoromethyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol;

(2R)-3-[[3-[3-(*N,N*-dimethylamino)phenoxy]phenyl][2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:

(2R)-3-[[[2-fluoro-4-(trifluoromethyl)phenyl]methyl][3-[[3-(trifluoromethoxy)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:

5 (2R)-3-[[[2-fluoro-4-(trifluoromethyl)phenyl]methyl][3-[[3-(trifluoromethyl)phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:

(2R)-3-[[[2-fluoro-4-(trifluoromethyl)phenyl]methyl][3-[[3,5-dimethylphenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:

10 (2R)-3-[[[2-fluoro-4-(trifluoromethyl)phenyl]methyl][3-[[3-(trifluoromethylthio)-phenyl]methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:

(2R)-3-[[[2-fluoro-4-(trifluoromethyl)phenyl]methyl][3-[[3,5-difluorophenyl]-methoxy]phenyl]amino]-1,1,1-trifluoro-2-propanol:

(2R)-3-[[[2-fluoro-4-(trifluoromethyl)phenyl]methyl][3-(cyclohexylmethoxy)-phenyl]amino]-1,1,1-trifluoro-2-propanol:

15 (2R)-3-[[3-(2-difluoromethoxy-4-pyridyloxy)phenyl][2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:

(2R)-3-[[3-(2-trifluoromethyl-4-pyridyloxy)phenyl][2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:

20 (2R)-3-[[3-(3-difluoromethoxyphenoxy)phenyl][2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol:

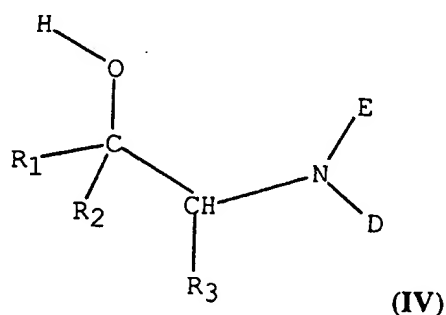
(2R)-3-[[[3-(3-trifluoromethylthio)phenoxy]phenyl][2-fluoro-4-(trifluoromethyl)-phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol: and

(2R)-3-[[3-(4-chloro-3-trifluoromethylphenoxy)phenyl][2-fluoro-4-(trifluoro-methyl)phenyl]methyl]amino]-1,1,1-trifluoro-2-propanol.

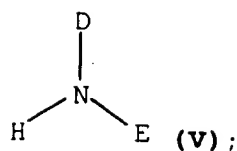
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31. A pharmaceutical composition comprising a compound of one of claims 1 through 30 together with a pharmaceutically acceptable carrier.

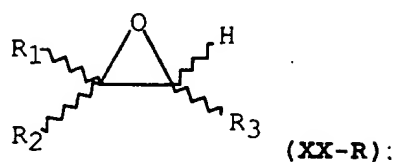
32. A method of treating coronary artery disease or other CETP-mediated disorders in a subject by administering a therapeutically effective amount of a compound of one of claims 1 through 30.
- 5 33. A method of preventing coronary artery disease or other CETP-mediated disorders in a subject by administering a therapeutically effective amount of a compound of one of claims 1 through 30.
34. A method of preventing cerebral vascular accident (CVA) in a subject by
10 administering a therapeutically effective amount of a compound of one of claims 1 through 30.
35. A method of preventing or treating dyslipidemia in a subject by administering a therapeutically effective amount of a compound of one of
15 claims 1 through 30.
36. A process for the preparation of compounds as recited in any one of claims 1 or 2 having the Formula (IV):



- 20 and pharmaceutically acceptable salts thereof, comprising the reaction of an amine of Formula (V):



with an epoxide of Formula (XX-R):



wherein:

- 5 R_1 is selected from the group consisting of haloalkyl and haloalkoxymethyl;

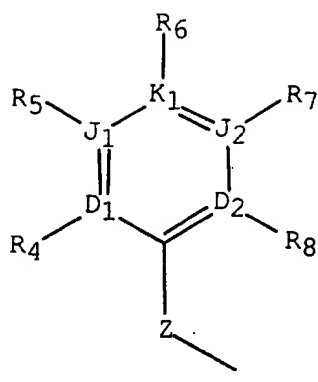
R_2 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, haloalkoxy, haloalkoxyalkyl, perhaloaryl, perhaloaralkyl, perhaloaryloxyalkyl, and heteroaryl;

- 10 R_3 is selected from the group consisting of hydrido, aryl, alkyl, alkenyl, haloalkyl, and haloalkoxyalkyl;

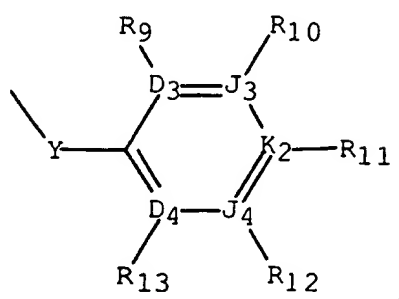
- D and E are independently selected from the group consisting of hydrido, A, and Q with the provisos that E and D are other than hydrido at the same time and A and Q are independently selected, when one of E and D is hydrido or when Y and Z are both single covalent bonds, from other than phenyl, 3-methylphenyl, 3-ethylphenyl, 2-methoxy-5-methylphenyl, 2-chlorophenyl, 3-chlorophenyl, and 3-bromophenyl;
- 15

A is the Formula:

300



Q is the Formula:



- D_1, D_2, J_1, J_2 and K_1 are independently selected from the group
 5 consisting of C, N, O, S and covalent bond with the provisos that no more
 than one of D_1, D_2, J_1, J_2 and K_1 is a covalent bond, no more than one of $D_1,$
 D_2, J_1, J_2 and K_1 is O, no more than one of D_1, D_2, J_1, J_2 and K_1 is S, one of
 D_1, D_2, J_1, J_2 and K_1 must be a covalent bond when two of D_1, D_2, J_1, J_2 and
 K_1 are O and S, and no more than four of D_1, D_2, J_1, J_2 and K_1 are N;
- 10 D_3, D_4, J_3, J_4 and K_2 are independently selected from the group
 consisting of C, N, O, S and covalent bond with the provisos that no more
 than one is a covalent bond, no more than one of D_3, D_4, J_3, J_4 and K_2 is O,

no more than one of D_3 , D_4 , J_3 , J_4 and K_2 is S, no more than two of D_3 , D_4 , J_3 , J_4 and K_2 are O and S, one of D_3 , D_4 , J_3 , J_4 and K_2 must be a covalent bond when two of D_3 , D_4 , J_3 , J_4 and K_2 are O and S, and no more than four of D_3 , D_4 , J_3 , J_4 and K_2 are N;

5 Y is selected from the group consisting of a covalent single bond,

$(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$ wherein j and k are integers independently selected from 0 and 1;

Z is selected from the group consisting of covalent single bond,

$(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$

10 wherein j and k are integers independently selected from 0 and 1;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group consisting of hydrido, halo, haloalkyl, and alkyl ;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the group consisting of hydrido, carboxy, heteroaralkylthio, heteroaralkoxy, cycloalkylamino, acylalkyl, acylalkoxy, aroylalkoxy, heterocyclyloxy, 15 aralkylaryl, aralkyl, aralkenyl, aralkynyl, heterocyclyl, perhaloaralkyl, aralkylsulfonyl, aralkylsulfonylalkyl, aralkylsulfinyl, aralkylsulfinylalkyl, halocycloalkyl, halocycloalkenyl, cycloalkylsulfinyl, cycloalkylsulfinylalkyl, cycloalkylsulfonyl, cycloalkylsulfonylalkyl, heteroaryl amino, N- 20 heteroaryl amino-N-alkyl amino, heteroaryl aminoalkyl, haloalkylthio, alkanoyloxy, alkoxy, alkoxyalkyl, haloalkoxyalkyl, heteroaralkoxy, cycloalkoxy, cycloalkenyloxy, cycloalkoxyalkyl, cycloalkylalkoxy, cycloalkenyloxyalkyl, cycloalkylenedioxy, halocycloalkoxy, halocycloalkoxyalkyl, halocycloalkenyloxy, halocycloalkenyloxyalkyl, 25 hydroxy, amino, thio, nitro, lower alkyl amino, alkylthio, alkylthioalkyl,

- arylamino, aralkylamino, arylthio, arylthioalkyl, heteroaralkoxyalkyl, alkylsulfinyl, alkylsulfinylalkyl, arylsulfinylalkyl, arylsulfonylalkyl, heteroarylsulfinylalkyl, heteroarylsulfonylalkyl, alkylsulfonyl, alkylsulfonylalkyl, haloalkylsulfinylalkyl, haloalkylsulfonylalkyl, 5 alkylsulfonamido, alkylaminosulfonyl, amidosulfonyl, monoalkyl amidosulfonyl, dialkyl amidosulfonyl, monoarylamidosulfonyl, arylsulfonamido, diarylamidosulfonyl, monoalkyl monoaryl amidosulfonyl, arylsulfinyl, arylsulfonyl, heteroarylthio, heteroarylsulfinyl, heteroarylsulfonyl, heterocyclylsulfonyl, heterocyclylthio, alkanoyl, alkenoyl, 10 aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, haloalkanoyl, alkyl, alkenyl, alkynyl, alkenyloxy, alkenyloxyalkyl, alkylenedioxy, haloalkylenedioxy, cycloalkyl, cycloalkylalkanoyl, cycloalkenyl, lower cycloalkylalkyl, lower cycloalkenylalkyl, halo, haloalkyl, haloalkenyl, haloalkoxy, hydroxyhaloalkyl, hydroxyaralkyl, hydroxyalkyl, hydroxyheteroaralkyl, 15 haloalkoxyalkyl, aryl, heteroaralkynyl, aryloxy, aralkoxy, aryloxyalkyl, saturated heterocyclyl, partially saturated heterocyclyl, heteroaryl, heteroaryloxy, heteroaryloxyalkyl, heteroaralkyl, arylalkenyl, heteroarylalkenyl, carboxyalkyl, carboalkoxy, alkoxycarboxamido, alkylamidocarbonylamido, arylamidocarbonylamido, carboalkoxyalkyl, 20 carboalkoxyalkenyl, carboaralkoxy, carboxamido, carboxamidoalkyl, cyano, carbohaloalkoxy, phosphono, phosphonoalkyl, diaralkoxyphosphono, and diaralkoxyphosphonoalkyl;

- R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , R_7 and R_8 , R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} are independently selected to form spacer 25 pairs wherein a spacer pair is taken together to form a linear moiety having from 3 through 6 contiguous atoms connecting the points of bonding of said spacer pair members to form a ring selected from the group consisting of a cycloalkenyl ring having 5 through 8 contiguous members, a partially saturated heterocyclyl ring having 5 through 8 contiguous members, a

heteroaryl ring having 5 through 6 contiguous members, and an aryl with the provisos that no more than one of the group consisting of spacer pairs R_4 and R_5 , R_5 and R_6 , R_6 and R_7 , and R_7 and R_8 , is used at the same time and that no more than one of the group consisting of spacer pairs R_9 and R_{10} , R_{10} and R_{11} , R_{11} and R_{12} , and R_{12} and R_{13} is used at the same time.

37. A process according to Claim 36 wherein the reaction is carried out at a temperature of from 0 °C to 100 °C.

38. A process according to Claim 37 wherein the reaction is carried out at a temperature of from 15 °C to 65 °C.

39. A process according to Claim 36 wherein the process further comprises a solvent selected from the group consisting of tetrahydrofuran, dioxane, methylene chloride, and acetonitrile.

40. A process according to Claim 36 wherein the process further comprises a transition metal salt catalyst selected from the group consisting of ytterbium, hafnium, scandium, neodymium, gadolinium, and zirconium salts.

41. A process according to Claim 40 wherein the transition metal salt is selected from the group consisting of ytterbium triflate, hafnium triflate, scandium triflate, neodymium triflate, gadolinium triflate, and zirconium triflate.

42. A process according to Claim 36 wherein the oxirane XX-R is selected from the group consisting of 2-trifluoromethyloxirane, 2-pentafluoroethyloxirane, 2-(1,1,2,2-tetrafluoroethoxymethyl)oxirane, 2-(difluorochloromethyl)oxirane, and 2-(trifluoromethoxymethyl)oxirane.

43. A process according to Claim 36 in which the oxirane has the (R)-chiral configuration at the R_1 and R_2 substituted carbon.

44. A process according to Claim 36, wherein:

5 D_1, D_2, J_1, J_2 and K_1 are each carbon:

D_3, D_4, J_3, J_4 and K_2 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that at least one of D_3, D_4, J_3, J_4 and K_2 is selected from the group consisting of O, S, and N,

wherein no more than one of D_3, D_4, J_3, J_4 and K_2 is a covalent bond, no

10 more than one of D_3, D_4, J_3, J_4 and K_2 is O, no more than one of $D_3, D_4, J_3,$

J_4 and K_2 is S, one of D_3, D_4, J_3, J_4 and K_2 must be a covalent bond when

two of D_3, D_4, J_3, J_4 and K_2 are O and S, and no more than four of $D_3, D_4,$

J_3, J_4 and K_2 are N.

15 45. A process according to Claim 36 wherein:

D_3, D_4, J_3, J_4 and K_2 are each carbon;

D_1, D_2, J_1, J_2 and K_1 are independently selected from the group consisting of C, N, O, S and covalent bond with the provisos that at least one of D_1, D_2, J_1, J_2 and K_1 is selected from the group consisting of O, S, and N,

20 wherein no more than one of D_1, D_2, J_1, J_2 and K_1 is a covalent bond, no

more than one of D_1, D_2, J_1, J_2 and K_1 is O, no more than one of $D_1, D_2, J_1,$

J_2 and K_1 is S, one of D_1, D_2, J_1, J_2 and K_1 must be a covalent bond when

two of D_1 , D_2 , J_1 , J_2 and K_1 are O and S, and no more than four of D_1 , D_2 ,

J_1 , J_2 and K_1 are N.

46. A process according to Claim 36 wherein D_1 , D_2 , J_1 , J_2 , K_1 , D_3 , D_4 , J_3 ,

J_4 and K_2 are each carbon.

5 47. A process according to Claim 36 wherein:

D_1 , D_2 , J_1 , J_2 , K_1 , D_3 , D_4 , J_3 , J_4 and K_2 are each carbon

Y is selected from the group consisting of a covalent single bond and

$(CH_2)_q$ wherein q is an integer selected from 1 and 2, and $(CH_2)_j-O-(CH_2)_k$

wherein j and k are integers independently selected from 0 and 1;

10 Z is a covalent single bond;

48. A process according to Claim 36 wherein:

D_1 , D_2 , J_1 , J_2 , K_1 , D_3 , D_4 , J_3 , J_4 and K_2 are each carbon

Y is selected from the group consisting of a covalent single bond and

C1-C2 alkylene;

15 Z is a covalent single bond;

R_4 , R_8 , R_9 , and R_{13} are independently selected from the group

consisting of hydrido and halo;

R_5 , R_6 , R_7 , R_{10} , R_{11} , and R_{12} are independently selected from the

group consisting of hydrido, alkyl, halo, haloalkyl, haloalkoxy, aryl,

20 alkylthio, arylamino, arylthio, aroyl, arylsulfonyl, aryloxy, aralkoxy,

heteroaryloxy, alkoxy, aralkyl, cycloalkoxy, cycloalkylalkoxy,

cycloalkylalkanoyl, heteroaryl, cycloalkyl, haloalkylthio, hydroxyhaloalkyl,

heteroaralkoxy, heterocyclyloxy, aralkylaryl, heteroaryloxyalkyl, heteroarylthio, and heteroarylsulfonyl.

49. A process according to any of Claims 44 through 48 wherein the reaction is carried out at a temperature of from 0 °C to 100 °C.
- 5 50. A process according to Claim 49 wherein the reaction is carried out at a temperature of from 15 °C to 65 °C.
51. A process according to any of Claims 44 through 48 wherein the process further comprises a solvent selected from the group consisting of tetrahydrofuran, dioxane, methylene chloride, and acetonitrile.
- 10 52. A process according to any of Claims 44 through 48 wherein the process further comprises a transition metal salt catalyst selected from the group consisting of ytterbium, hafnium, scandium, neodymium, gadolinium, and zirconium salts.
53. A process according to Claim 52 in which the transition metal salt is
15 selected from the group consisting of ytterbium triflate, hafnium triflate, scandium triflate, neodymium triflate, gadolinium triflate, and zirconium triflate.
54. A process according to any of Claims 44 through 48 wherein the oxirane XX-R is selected from the group consisting of 2-trifluoromethyloxirane,
20 2-pentafluoroethyloxirane, 2-(1,1,2,2-tetrafluoroethoxymethyl)oxirane, 2-(difluorochloromethyl)oxirane, and 2-(trifluoromethoxymethyl)oxirane.
55. A process according to Claim 54 in which the oxirane has the (R)-chiral configuration at the R₁ and R₂ substituted carbon.
56. A process according to any of Claims 44 through 48 wherein the process
25 further comprises a:

- (a) Temperature of from 0 °C to 100 °C;
- (b) Non-protic solvent;
- (c) Transition metal salt selected from the group consisting of
ytterbium, hafnium, scandium, neodymium, gadolinium, and
zirconium salts.

5

57. A process according to Claim 56 wherein the reaction is carried out at a temperature of from 15 °C to 65 °C.

58. A process according to Claim 56 wherein the solvent is selected from the group consisting of tetrahydrofuran, dioxane, methylene chloride, and acetonitrile.

10

59. A process according to Claim 56 wherein the transition metal salt is selected from the group consisting of ytterbium triflate, hafnium triflate, scandium triflate, neodymium triflate, gadolinium triflate, and zirconium triflate.

60. A process according to Claim 56 wherein the oxirane XX-R is selected from the group consisting of 2-trifluoromethyloxirane, 2-pentafluoroethyloxirane, 2-(1,1,2,2-tetrafluoroethoxymethyl)oxirane, 2-(difluorochloromethyl)oxirane, and 2-(trifluoromethoxymethyl)oxirane.

15

61. A process according to Claim 60 wherein the oxirane has the (R)-chiral configuration at the R₁ and R₂ substituted carbon.

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 99/22120

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C217/90 C07C217/84 C07D263/06 C07D251/16 A61K31/135
A61K31/33 C07C217/82 C07C217/86 C07C239/20

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|----------|--|-----------------------|
| X | DUNN C ET AL: "THE SYNTHESIS OF FLUORINE-CONTAINING PTERINS" TETRAHEDRON, NL, ELSEVIER SCIENCE PUBLISHERS, AMSTERDAM, vol. 52, no. 40, page 13017-13026 XP002063653 ISSN: 0040-4020 page 13024, line 4 - line 16 | 1 |
| A | EP 0 801 060 A (PFIZER) 15 October 1997 (1997-10-15) cited in the application abstract | 1, 32-35 |
| A | GB 2 305 665 A (MERCK & CO INC) 16 April 1997 (1997-04-16) cited in the application abstract; claims 1-17 | 1, 32-35 |
| -/- | | |

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "3" document member of the same patent family

Date of the actual completion of the international search

11 January 2000

Date of mailing of the international search report

21/01/2000

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 99/22120

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|----------|---|-----------------------|
| A | US 2 700 686 A (JOSEPH B. DICKEY ET AL.) 25 January 1955 (1955-01-25) cited in the application abstract --- | 1 |
| A | EP 0 818 197 A (BAYER AG) 14 January 1998 (1998-01-14) abstract --- | 1, 32-35 |
| A | KATAGIRI T ET AL: "Intramolecular SN2 reaction at alpha-carbon of trifluoromethyl group: preparation of optically active 2-trifluoromethylaziridine" TETRAHEDRON: ASYMMETRY, NL, ELSEVIER SCIENCE PUBLISHERS, AMSTERDAM. vol. 8, no. 17, page 2933-2937 XP004090383 ISSN: 0957-4166 page 2936, paragraph 3 ----- | 1 |

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 99/ 22120

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 32-35
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 32-35
are directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☒ Claims Nos.: 1-31, 36-61
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report
covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
restricted to the invention first mentioned in the claims: it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/SA/ 210

Continuation of Box I.2

Claims Nos.: 1-31, 36-61

Present claims 1-31, 36-61 relate to an extremely large number of possible compounds/methods. In fact, the claims contain so many options, variables, and provisos that a lack of clarity (and/or conciseness) within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible. Consequently, the search has been carried out for those parts of the application which do appear to be clear (and/or concise), namely according to formula (I) of claim 1, wherein:

X is O

R1 = CF₃

R16 = free site

n = 1

R3 = R14 = R15 = Hydrogen

Y = C, O, bond

D1 = D3 = D4 = J3 = J4 = K2 = C

the other substituent on the Nitrogen atom is -Ar, CH₂-Ar, Het, -CH₂-Het cycle with free sites.

It is stressed that this scope covers the majority of the examples.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

In International Application No

PCT/US 99/22120

| Patent document cited in search report | Publication date | Patent family member(s) | Publication date |
|---|---------------------|--|--|
| EP 0801060 A | 15-10-1997 | CA 2201988 A JP 10036348 A US 5843972 A | 09-10-1997 10-02-1998 01-12-1998 |
| GB 2305665 A | 16-04-1997 | US 5714506 A | 03-02-1998 |
| US 2700686 A | 25-01-1955 | NONE | |
| EP 0818197 A | 14-01-1998 | DE 19627431 A BG 101748 A BR 9703890 A CA 2209825 A CN 1174196 A CZ 9702144 A HR 970333 A HU 9701157 A JP 10167967 A NO 973143 A PL 320953 A SG 46781 A SK 92597 A US 5932587 A | 15-01-1998 30-04-1998 03-11-1998 08-01-1998 25-02-1998 14-01-1998 30-04-1998 30-03-1998 23-06-1998 09-01-1998 19-01-1998 20-02-1998 06-05-1998 03-08-1999 |